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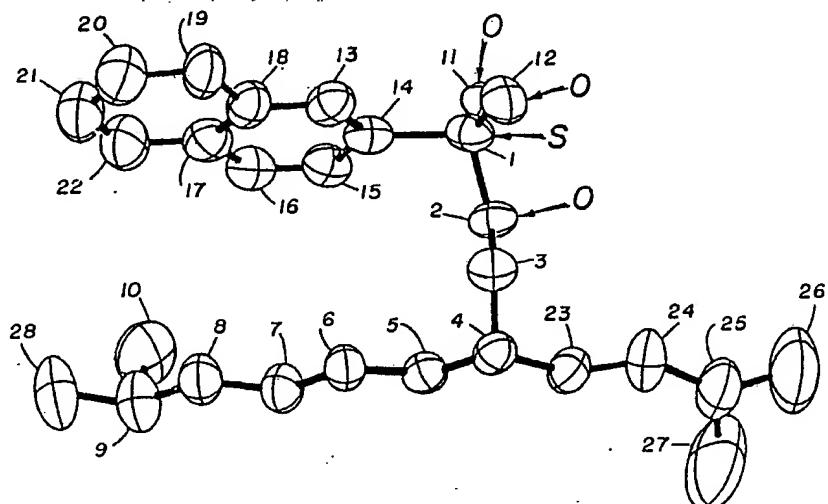
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(54) Title: DETERGENT RANGE ALDEHYDE AND ALCOHOL MIXTURES AND DERIVATIVES, AND PROCESS THEREFOR



(57) Abstract

Novel, liquid mixtures of isomeric aldehydes and alcohols are described in the C₁₁-C₁₆ carbon range, the compounds being characterized by a main carbon branched at the position and moderate additional branching in most isomers; the aldehyde mixtures are prepared by an economic route from olefins involving oxo and aldol reaction with the reaction conducted in such a way as to give a high percentage of aldolable product, and preferably with a base catalyzed aldol reaction conducted under conditions to make high conversions attainable. The aldehyde mixtures can be hydrogenated to alcohols and converted to novel ethoxylates of sulfate compositions suitable for use as biodegradable detergents; or hydrogenated and oxidized to novel carboxylic acid compositions also suitable for detergent use.

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DETERGENT RANGE ALDEHYDE AND ALCOHOL MIXTURES
AND DERIVATIVES, AND PROCESS THEREFOR

5 The present invention is concerned with a process for preparing detergent hydrophobes from olefin feedstock, and with novel detergent range aldehyde, alcohol and acid hydrophobes.

BACKGROUND OF THE INVENTION

10 The principal commercial surfactants in use today are linear alkylbenzene sulfonates and linear alcohol ethoxylates. The hydrophobic portion of both surfactants is a linear alkyl chain of between eleven and eighteen carbon atoms. These surfactants had been preceded by synthetic detergents which contained highly branched groups as a hydrophobic portion. The 15 change to the currently employed linear groups, which occurred in the 1960's, was prompted by concern over the slow biodegradation characteristics of the branched hydrophobes. The perceived need for linearity led to development of particular approaches 20 to hydrophobe preparation. The linear alkyl benzene sulfonates are based upon linear olefins derived from paraffins, which in turn are obtained through molecular sieve separations from paraffin mixtures. The linear alcohol based detergents are produced by 25 way of ethylene oligomerization, followed by processes to manipulate the broad range of olefins obtained into the desired molecular weight range. The processing involved in such approaches adds considerably to energy and facilities usage and consequently to



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product cost. When olefins of the requisite carbon atom number have been obtained, they can be hydroformylated to produce aldehydes, generally with one more carbon atom than the olefin, which can be 5 hydrogenated to an alcohol.

5 An alternate method for generating longer chain alcohols from short chain olefins is via a sequence involving hydroformylation (or oxo reaction) followed by aldol condensation and hydrogenation.

10 Thus 2-ethylhexanol is prepared on a very large scale by (a) hydroformylating propylene to a mixture of n-butanal and isobutanal, (b) separating the mixtures of aldehydes (c) aldol reaction of n-butanal to 2-ethylhexenal and (d) hydrogenation of 2-ethylhexenal 15 to 2-ethylhexanol. While this approach is well-recognized to be cost effective for generation of medium chain alcohols, it has not heretofore been shown to be an economical method for generation of longer chain alcohols. Among patents teaching 20 conversion of aldehydes to higher aldehydes by the well known aldol reaction is U.S. patent 2,852,563.

Medium chain length olefins are usually derived from dimerization or oligomerization of ethylene or propylene. Among dimerization processes is 25 the Dimersol® dimerization process for dimerizing olefins using a nickel coordination complex and an aluminum alkyl as catalyst. The process can convert propylene to hexenes with selectivity in excess of 85%. The hexenes can be converted by oxo reaction to 30 aldehydes and then alcohols, producing heptanols. Processes are also known for dimerizing propylene with trialkylaluminum metals to 4-methyl-1-pentene, see Industrial Organic Chemistry, Klaus Weissermel and Hans-Jurgen Arpe; English translation by Alexander 35 Muller (Verlag Chemie, Weinheim, New York, 1978), pp



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75-77. Also oxo reactions of certain branched olefins have been studied; see M. Johnson, . Chem Soc. 1963, 4859; Piacenti et al, J. Chem. Soc., 1966, 488; and Vysokinskii et al, J. Applied Chemistry of USSR, 1972, 5 Vol. 45, pp. 1352-1355. Also oxo reactions of certain non-terminal octenes have been reported to give less than 60% of the straight chain aldehyde isomers, see Kummer et al, Homogeneous Catalysis-II, pages 19 to 26, Advances in Chemistry Series 132 (Edited by Denis 10 Forster and James F. Roth), American Chemical Society, Washington D.C., 1974.

Reactions of the type described characteristically produce mixtures of products, often with extensive branching. Therefore, in order to 15 control the branching, or to eliminate unreactive components, it has been common practice to employ distillation at intermediate stages to remove some of the isomers.

SUMMARY OF THE INVENTION

20 The invention is concerned with a process in which olefins selected from those with 3 to 7 carbon atoms are converted to aldehydes with 4 to 8 carbon atoms, which are then subjected to aldol condensation to obtain aldol products in relatively high yield. 25 The aldehydes can be hydrogenated to alcohols having useful properties and derivative uses, with especial interest in conversion of hexenes to heptanals and to C₁₄ alcohols for use in detergents. The aldehydes can also be converted to other compounds useful as 30 detergents or for other purposes. The invention is particularly concerned with the foregoing process in which the olefins have 5 to 7 carbon atoms and are converted to aldehydes with 6 to 8 carbon atoms, and then to enals and saturated aldehydes with 12 to 16 35 carbon atoms, in the detergent hydrophobe range. Such



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processes and the aldehyde, alcohol and detergent derivative products thereof are particularly exemplified herein by processes leading to, and products mainly composed of, compounds with 14 carbon atoms. The conditions of the exemplary process are generally applicable to those employing other reactants in the stated carbon atom ranges.

The present invention particularly concerns a hydrophobe aldehyde or alcohol mixture composed almost entirely of C_{14} aldehydes or alcohols with a structure branched at the 2-position, with the alcohols being generally 2-pentylnonanols, i. e. having a five-carbon alkyl group substituted on the 2-position of a nine-carbon alkanol; with most of the isomers in the mixture having moderate additional branching, primarily methyl groups. The mixture is liquid at ambient temperatures and the hydrophobe groups of the alcohols are such as to make the mixture useful for formation of very effective detergents.

The alcohol structures are further characterized by the absence of quaternary carbon or other structures strongly resistant to biodegradation, while generally having vicinal branching within limited ranges, and are still suitably biodegradable. The invention is further concerned with other aldehydes and alcohol mixtures of such 2-branched structure, but with 11 to 16 carbon atoms., e.g. ethyl-nonanol and hexyldecanols.

The invention is further directed to processes for converting olefins to described C_{14} aldehydes and alcohols, involving dimerization, oxo, aldol and hydrogenation processes based on propylene. In particular, a process involves conducting an oxo reaction with a hexene mixture comprised mainly of methyl pentenes, as produced by dimerization of



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propylene, and obtaining a heptanal product in which carbonylation has occurred primarily on terminal carbon so that upwards of 75% of the heptanal isomers are unbranched at the 2-position and capable of 5 reacting directly in aldol reactions under basic conditions; and conducting an aldol reaction of the heptanal product to obtain C₁₄ aldehydes which are then hydrogenated to C₁₄ alcohols. The invention further includes use of cobalt catalyst in the oxo 10 procedure under conditions to produce high percentages of desired aldolable product, and utilization of a co-solvent and suitable aldol conditions to obtain the C₁₄ product. Fortuitously, it has been found that in the hydroformylation of isomeric hexenes, the 15 formylation of the prominent 2-methyl-2-pentene isomer occurs primarily on a terminal bond, with the percentage of such terminal formylation being much higher than that generally characteristic of hydroformylation of internal olefins; and a high 20 percentage of aldolable product is obtained from terminal formylation of isomeric hexenes. The present invention involves an efficient process for converting propylene dimers to detergent hydrophobe alcohols in high overall yields such as 70% or better, and the 25 alcohols are suitable for conversion to detergents having detergent and biodegradation properties comparable to those of the common commercial detergents. Moreover, the present process provides a much more efficient and economical route to detergent 30 hydrophobes from olefin feedstock than that provided by the routes prominent in present commercial use. This route permits detergent alcohols to be built up from low cost propylene as compared to present 35 commercial processes based on ethylene. In the processes of the invention, the immediate precursor of



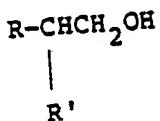
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an alcohol is an aldehyde, and the aldehydes can also be converted to other useful compounds.

DETAILED DESCRIPTION OF THE INVENTION

The C₁₄ alcohols of the present invention

5 can be represented:



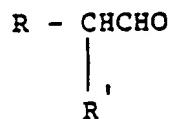
in which R is an alkyl group with seven carbon atoms and R' is an alkyl group with five carbon atoms. In a high percentage of the alcohols in the mixture, 10 generally more than 80-85% or so, R' is selected from n-pentyl, 3-methylbutyl and 1-methylbutyl, and R is selected from n-heptyl, 3-methylhexyl, 5-methylhexyl, 2-methylhexyl and 2-ethylpentyl groups. While there will be some variance in the alcohol mixture with 15 variation in preparation conditions, generally close to 50% or so of the alcohol mixture will be composed of 2-(3-methylbutyl)-5-methyl-octanol, 2-(1-methylbutyl) 5-methyloctanol and 2-pentyl-5-methyloctanol. Considering the type of branching involved, close to 20 50% or so by weight of the alcohols will have branching not only at the 2-position, but with each branch having an additional methyl branch, i.e. with R' and R in the above formula being selected from methylbutyl and methylhexyl groups respectively. The 25 alcohols are also characterized as generally having a limited amount of vicinal substitution, i.e. substitution or branching on adjacent carbon atoms, and little or virtually no di-substituted carbon chain atoms, i.e. quaternary carbon atoms. Significant 30 amounts of alcohols with quaternary carbon atoms have not been found in product analyses, and the types of



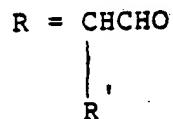
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reactants and reactions used in product preparations make it very unlikely that significant amounts of such alcohols would be found in unidentified portions of products. The lack of quaternary carbon in the 5 alcohols of the mixture is fortunate in that such carbon is ordinarily resistant to biodegradation.

The described alcohols are obtained by hydrogenation of aldehydes of the same structure, and such aldehydes are similarly useful for their 10 hydrophobe character and convertability to useful detergent or other derivatives. Such aldehydes are represented by the formula:



or when still in the enal form:



where the double bond is to one of the carbons of the 15 R group. In such formulae, R and R' have the same meanings as described with respect to the alcohols, except that R in the enal is now alkylidine. For alcohols, enals and saturated aldehydes of twelve 20 carbon atoms, R has six carbon atoms, and R' has four; and for sixteen carbon atoms R has eight carbon atoms and R' has six. R and R' will similarly vary for other structures having 11 to 16 carbon atoms, R from 6 to 8 carbon atoms, and R' from 2 to 6 carbon atoms. As with the C₁₄ alcohols, the C₁₁, C₁₂, and C₁₆ 25 aldehydes and alcohols will often have branching not



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only at the 2-position, but with each branch having an additional methyl or ethyl branch.

A useful group of the unsaturated aldehydes of the present invention can also be represented:



in which R plus R' total 8 to 13 carbon atoms and in which often in a high percentage (75%) of the unsaturated aldehyde, alkyl branches R and R' can be branched also with branching being limited to methyl

10 or ethyl groups.

With further regard to the particular C₁₄ alcohol mixture, approximately 40 to 60% by weight will generally be the 2-(3-methylbuty)-5-methyloctanol, 2-(1-methylbutyl)-5-methyloctanol and 2-pentyl 15 -5-methyloctanol; and at least about 65% and possibly 65 to 80% or so by weight will be comprised of these alcohols, along with 2-(3-methylbutyl)-7-methyloctanol, 2-pentynonanol, 2-pentyl-7-methyloctanol and 2-(3-methylbuty)-nonanol. In 20 addition to the indicated ranges of these alcohols, various other C₁₄ isomers will be present in small percentages, particularly those other alcohols listed in Table 4 herein, in amounts by weight usually approximating those in the Table and generally no more 25 than 5% each and usually in ranges up to 1 to 2% or so, and with the balance of the C₁₄ alcohols being generally composed of the other alcohols in Table 3 herein, which will generally be present only in small percentages of each isomer. It will be noted that 30 those isomers present only in small amount, with no amount reported in Table 4, but included in Table 3, tend to be more highly branched, i.e. multi-branched,



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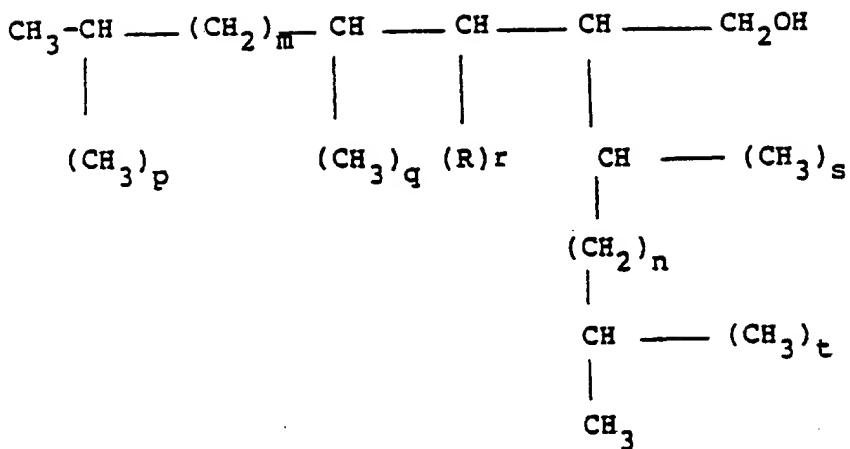
than the other isomers, although most branches are still methyl, or occasionally ethyl groups. Taking these isomers into consideration, the amount of vicinal branched material may range as high as 20 to 5 35% or so by weight of the C₁₄ alcohols. However, the mixtures of alcohol isomers are biodegradable as indicated by tests reported herein. One of the alcohols which can be present in fairly good proportion, such as 10 to 20% or so, 10 2-(1-methyl-butyl)-5-methyloctanal is a vicinal branched alcohol. The alcohols in the mixture are characterized by a fair degree of branching, although the alcohols present in large amount tend to be less branched than some of the minor components. Thus the 15 18 or so alcohols constituting about 90-95% or so by weight of the alcohols have an average of slightly more than 2 branches, about 2.2 to 2.4 branches, while the other 5 to 10% of the alcohols may have upwards of above 3.5 branches, such as 3.75-3.8 branches.

20 In addition to the novelty of the C₁₄ alcohol mixtures of the present invention, the particular alcohols are also new except for the 2-pentylnonanol. Thus the other 27 alcohols named in Table 3 are new compounds suitable for formation of 25 nonionic detergents.

The alcohols comprising most of the present alcohol mixtures produced from branched hexenes can also be represented by the formula:



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wherein:

R = methyl or ethyl;

p, q and r = 0 or 1; but only one of p, q

5 and r can be 1;

m = 3, when p, q and r = 0;

m = 2, when p or q = 1;

m = 2, when r = 1 and R = methyl;

m = 1, when r = 1 and R = ethyl;

10 n = 2, when s and t = 0;

n = 1, when s or t = 1;

s and t = 0 or 1, but only one of s and t can be 1.

15 The alcohol mixtures, and the individual components thereof, are liquids at ambient temperatures. This is advantageous in that the alcohols can be more readily transferred from one vessel to another, or moved by pumping through conduits, etc., than is the case with solid alcohols.

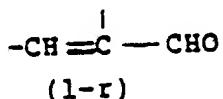
20 Also the liquid form is more convenient for mixing with reactants and solvents for conversion to detergent compounds or other useful products as contemplated. It happens that C_{14} alcohols with straight chains, or with considerably less branching

25 than the present alcohols, are generally solids. Thus

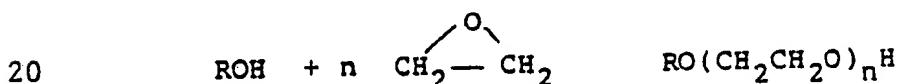


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5 detergent alcohols obtained by way of ethylene oligomerization, such as 75% to 80% normal primary C_{12-15} alcohols and 75% normal C_{14} primary alcohols, consisting of the designated percentages of normal alcohols and the balance of isomeric 2-alkyl (primarily 2-methyl) primary alcohols, are solid materials. Mixtures of aldehydes can also be represented by the above formula, with the CH_2OH group replaced by CHO . For the corresponding enal structure, there is 10 unsaturation at the 2-position:



15 The detergent alcohols of the present invention are particularly suitable for conversion to nonionic detergents of good deterutive and biodegradation properties. The largest volume use of detergent alcohols is as nonionic surfactants. The polyethoxylate surfactants are particularly important, and can be generated from the present alcohols by base catalyzed reaction between ethylene oxide and alcohol:



Typical values of n are in the range of 6 to 12, and R represents the alkyl portion of the alcohol, which for the present alcohols is generally a 14-carbon alkyl group. Ethoxylates prepared from the present alcohols 25 were tested for detergency and found to be comparable to a leading commercial nonionic detergent under a variety of washing conditions. The commercial detergent is that prepared from C_{12-15} alcohols of



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75-80% linearity, Neodol[®] 25.

In addition to deterutive properties, another concern with surfactants has been biodegradability. The first major synthetic detergents were so-called alkylbenzene sulfonates, with an alkyl hydrophobe group derived from propylene tetramer. The propylene tetramer was produced by acid catalyzed polymerization and hence was highly branched, including extensive quaternary branching. This gave rise to a product with rather poor biodegradation properties, which led to extended life for surfactant properties in rivers and lakes. Public concern over the asthetic impact (foam) and possible toxicity of long lasting surfactants led to a voluntary change over to predominantly linear hydrophobes. Prominent among current surfactants are linear alkylbenzene sulfonate and the linear alcohol ethoxylates discussed above. The linear alkylbenzene sulfonate (LAS) involves a linear alkyl group substituted on a benzene ring, generally at one of the secondary carbon atoms of the alkyl group. LAS has been found to be a suitably biodegradable detergent, although being degraded somewhat more slowly in standard tests than the substantially linear alcohol ethoxylates. The ethoxylates produced from the alcohols of the present invention are comparable in biodegradation to LAS, and therefore suitable in this regard. It is fortunate to find that the present alcohols are biodegradable, despite the presence of multiple branching. The substantially linear alcohol ethoxylates, such as Neodol[®] 25, are known to be biodegradable, but those alcohols include substantial percentages of linear alcohols, and the branching is ordinarily only a single methyl or other lower alkyl group. In contrast, most of the alcohols in the present mixture



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have more than two alkyl branches. It is advantageous to be able to produce the present detergent alcohols, of properties comparable to Neodol [®] alcohols, in a more efficient process and from a less expensive 5 feedstock, i.e. from propylene rather than from ethylene.

The enals and aldehydes produced in the present process are also suitable for conversion to other useful compositions. Thus, in accord with the 10 discovery of one of the present applicants, olefins can be converted to useful amine compounds by reactions involving hydroformylation and aldol reactions, followed by reaction with amines to obtain substituted amines. The reaction can be effected at 15 the enal stage, or with the saturated aldehydes. Thus tetradec-2-enals or tetradecanals are reacted with a secondary amine under reductive conditions to obtain tetradecyl amines, e.g. dimethyl amine gives N,N-dimethyltetradecylamines. Use of a primary amine 20 gives di(tetradecylamines), e.g. with mono-methyl amine, di-(tetradecyl)methylamines are produced. The amines with one or two long chain hydrophobe groups have detergent properties, and can be oxidized to amine oxides which also have detergent properties, or 25 quaternized to ammonium salts which have interesting germicidal and other properties. Thus the present invention provides aldehydes which can be utilized for production of amines or ammonium compounds with useful properties. The aldehydes can also be converted to the 30 corresponding saturated or unsaturated acids in accord with procedures described herein. While some branched-chain acids in the detergent range have an objectionable odor, the C₁₁ to C₁₆ derivatives of the aldehydes described herein are expected to have 35 suitably low odor in view of the type of branching



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involved.

Hexenes, as produced by dimerization of propylene with transition metal catalysts, as in the Dimersol® dimerization process, are characterized by 5 being composed mainly of internal olefins, and a linear content which has a range from about 20% up to 32% or so. The main isomer present is a 2-methyl-2-pentene, along with other 2- and 4-methyl pentenes and around 6% 2,3-dimethyl-2-butene.

10 As indicated above, the linear content of propylene dimers is fairly low, being around 20% to 30% or so. At times there may be advantage in separating the linear components prior to conducting the present process. Separation can be effected by 15 use of a molecular sieve or other suitable procedure. However, it has been found that the entire hexenes portion of the propylene dimerization product can be utilized as feedstock for the present oxo-aldol process. The branched isomers are typified by 20 2-methyl-2-pentene which, when subjected to oxo reaction with cobalt catalyst, has been found to be very selectively converted to 3-methyl and 5-methylhexanals. Fortunately it has been found that cobalt catalyst, in contrast to rhodium, has the 25 effect of isomerizing the internal olefins so that the aldehyde group is predominantly on the end of the chain. It is very important to the use of propylene dimer in the present process, that the oxo product is predominantly a non 2-branched alkanal., i.e. there is 30 no substituent in the 2-position. Such aldehydes, which will directly react in base-catalyzed aldol reactions are sometimes referred to herein as "aldolable" aldehydes. As discussed herein, aldehydes with substituents in the 2-position do not readily 35 undergo base-catalyzed aldol reactions. Thus if the



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internal hexenes were converted largely to such unreactive aldehydes, it would be very difficult to effect self-condensation of such aldehydes to a useful extent, and the use of propylene dimers in the present 5 oxo-aldol process would be impractical. However, as discussed herein, the oxo process with cobalt catalyst converts the hexenes largely, e.g. 75 to 80% or so, to aldehydes which will react in the aldol reaction, making hexenes, obtained from propylene dimerization 10 very suitable as a feedstock for producing detergent range alcohols in accord with the present invention. This result is surprising in view of the fact that oxo reactions of certain non-terminal octenes are reported to give less than 60% of straight chain aldehyde 15 isomers. It appears that the methyl substituent has some influence in directing the oxo reaction to obtain a great predominance of aldehydes with no substituent in the 2-position. There will similarly be advantage in using branched pentenes and heptenes in the present 20 process, including those with no more than 20 to 30% linear content, even including those with mainly internal unsaturation, and obtain similarly good results in the oxo and aldol reactions taught herein.

Particular branched hexene isomers are 25 converted to non 2-branched alkanals with very high selectivity, with 2-methyl-pentene-1 selectivity of better than 90% to such aldehydes being obtainable. The mixture of both branched and linear hexenes from propylene dimerization can be converted to such 30 aldolable aldehydes with selectivity such as about 79% or so. In contrast, the selectivity to such aldehydes from the linear hexenes may be only 60% or so. Thus, surprisingly it is found that higher selectivity to aldehydes desirable for the aldol reaction can be 35 obtained by using the crude hexenes mixture for the



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oxo reaction, rather than only the linear hexenes. Depending upon relative value and availability of the linear and branched hexenes, one might find advantage in using only the branched hexenes in the present 5 process because of the high selectivity in the oxo process to aldehydes with no 2-branching suitable for aldol reaction.

The linear hexenes can be separated from the crude dimerization product by use of a molecular sieve 10 or other suitable procedure, and the linear hexenes alone then subjected to an oxo reaction to obtain heptanal products, with the linear content of the heptanals ranging up to 75% or more. Branched isomers which may be present include 2-methylhexanal, and 15 2-ethylpentanal. The oxo product mixture can be reacted in an aldol reaction, employing aldol conditions as described herein, to produce aldol products. Under the aldol conditions employed, the n-heptanal reacts with itself at a rate about 10-15 20 times faster than it reacts with 2-methyl-hexanal or other 2-substituted aldehydes. Thus, when the C₇ aldehydes are produced from linear hexenes, a product compound predominantly of product from 25 self-condensation of n-heptanal can be obtained by carrying out an aldol reaction of the oxo reaction product, with some control over the amount of product from 2-substituted aldehydes by controlling the amounts of conversion which is permitted. It may be desirable to have better than 50 to 60% completion for 30 efficient use of feed stock, and conversion of 80% or higher may at times be desirable. The reaction can be run to achieve 95% or better conversion of the n-aldehyde to aldol product, while only about 1/4 to 1/3 or so of the 2-substituted aldehydes are usually 35 converted to aldol product by a cross-aldol reaction.



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With use of appropriate control, aldol product with about 80% to say 95% from self-condensation of n-heptanal can be obtained, for example at least 85% from self-condensation, with no more than 15% of 5 cross-aldol product. Depending upon the properties desired in the product, the degree of branching can be controlled to a considerable extent by the present process. It happens that some known detergent range 10 alcohols have a fair degree of branching and still have satisfactory biodegradability. Regardless of the desired degree of branching, there is advantage in being able to carry out an aldol reaction on the oxo 15 product of the hexenes, without need for separating branched aldehyde isomers, and obtain a useful product, particularly considering the low cost nature of the feed stock and process. With a proper heptanal mixture, the process can be used to obtain a product composed of about 85% 2-pentylnonanol and 15% 2-pentyl-4-methyl-octanol.

20 The oxo process can generally be utilized to achieve 90-95% yields of aldehydes. With linear hexenes as reactant, selectivity to aldehydes without 2-substituents, i. e. α -aldehydealkanes, can be as high as 60 to 65%, but in large scale operation will 25 possibly range from 50 to 65%.

The present invention employs an oxo reaction of substantially branched hexenes to obtain a mixture of aldehydes, which is then subjected to an aldol reaction. The oxo reaction involves contacting 30 hexenes with hydrogen and carbon monoxide and hydroformylation catalyst under hydroformylation conditions suited to obtaining a high proportion of terminal formylation of the olefin feed. It is desirable to have the resulting aldehyde constitute 75 35 to 80% or more of the aldehyde product.



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It is important that the hydroformylation of the mixed hexenes give a relatively high ratio of aldehydes without 2-branching, as this contributes to the feasibility of using the aldehyde mixture for an 5 aldol reaction to obtain a good yield of aldol product. The use of moderate temperatures in the hydroformylation contributes to forming aldehydes without 2-branching, but reaction rate improves with 10 temperature. Thus temperatures sufficient to produce an appreciable reaction rate, ranging from 80° to 100° C. or so can be used, and temperatures on up to 125-140° C. can be employed to obtain better reaction rates. Still higher temperatures up to 150° C. or 15 higher can be used. To some extent high catalyst concentrations can be employed to obtain reaction 20 rates, even at relatively low temperatures. Cobalt catalyst is especially suited to obtain the desired high proportion of aldehyde with no 2-branches. Unmodified cobalt carbonyl catalyst can conveniently 25 be used. Such catalyst conventionally designated as dicobalt octacarbonyl, can be provided or employed in many forms known to be useful as a hydroformylation catalyst, although it may be necessary to exercise some choice to provide catalyst best suited to obtaining a high proportion of aldehyde with product 30 suitable for direct base-catalyzed aldol reaction. The above-referred to S.N. 272,587 disclosed conditions as above for hydroformylation of mixed linear butenes, stating that moderate temperatures in the hydroformylation contribute to obtaining about a 3:1 35 mixture of normal to branched aldehydes. The conditions can be used for hydroformylation of olefins with 3 to 7, or more narrowly, 5 to 7 carbon atoms. The S. N. 272,587 application further described as exemplary a process in which mixed butenes are



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converted to a ten-carbon plasticizer alcohol comprised of at least about 80-90% 2-propylheptanol by an oxo reaction of the butenes to obtain amyl aldehydes with at least about 66% n-pentaldehyde content, followed by an aldol reaction of the aldehydes under conditions to cause substantially all of the n-pentaldehyde to react but with incomplete conversion of branched aldehydes, and then hydrogenating to produce alcohols in which the 10 ten-carbon alcohols are comprised of at least 80-90% 2-propylheptanol. Under the aldol conditions employed the 2-methylbutanal present does not readily condense with itself, and condenses at a comparatively slow rate with the n-pentanal, so that the 15 2-propyl-4-methylhexanol content (resulting from the so-called cross aldol of n-pentanal with 2-methylbutanal) in the resulting alcohol is held to no more than about 15-20%, often 12% or less. Under the aldol conditions employed, the n-pentanal reacts 20 with itself to form aldol product at a rate about 15 times greater than it reacts with 2-methylbutanal. The aldol reaction is permitted to go to 80% or so completion so that if about 25% of the aldehyde supplied is 2-methylbutanal, about 3/4 of it will 25 remain unreacted, and about 88% of the aldol product will be that from self-condensation of n-pentanal. The conversion of n-pentanal to aldol product will be very high and desirably nearly complete, such as upwards of 90 or 95%. There will be some variation 30 with conditions and isomer content of the aldehydes utilized, but it was contemplated to obtain aldol product with about 80% to about 95% being from the self-condensation of n-pentanal, and preferably at least 85% from self-condensation with no more than 15% 35 of 2-propyl-4-methylhexanal being produced. In the



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process, the aldol intermediate, 2-propyl-3-hydroxyl-heptanal, will ordinarily be dehydrated in the aldol procedure to 2-propylheptenal. Under some conditions the immediate aldol product can be 5 isolated, but ordinarily under the temperature conditions employed the 2-propyl-2-heptenal is produced. In such procedure, it was desirable to have the n-pentanal to branched aldehyde ratio at least about 2.0:1, representing at least about 66.7% 10 n-pentanal content. Aldehydes with 70-75% normal content, or even higher normal contents are desirable to the extent available from oxo reactions, possibly up to 85%, and will be useful for the aldol stage.

The oxo stage of the reaction can be 15 conducted under the usual conditions pertaining to cobalt catalyzed hydroformylation reactions with attention to the temperature conditions as described above. Usual pressure conditions apply, such as 500-4000 or up to 5000 psi (3447.5-27,580 on up to 20 34,475 kilopascals) total pressure, with most of the pressure being from the carbon monoxide and hydrogen supplied. The carbon monoxide and hydrogen are conveniently used in 1:1 ratio and obtained from usual synthesis gas sources, but other ratios can be 25 employed in keeping with known hydroformylation practice. The reaction can be carried to the desired stage of completion in 1 to 3 hours or so on a batch basis, temperature, pressure and catalyst concentration.

30 The reaction can be conveniently conducted either without a solvent or with solvents and, employing concentrations customary for homogeneous catalyst reactions, such as 2 to 10 molar or greater concentrations of the hexenes (or butenes or other 35 olefins) in a solvent, e.g. hydrocarbon solvents such



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as toluene, and 0.1% to 1% by weight, based on cobalt, of catalyst.

The present invention is particularly concerned with preparing detergent range alcohols and 5 aldehydes from propylene feedstock. The detergent range alcohols are somewhat higher in carbon number than plasticizer alcohols, often having 14 carbon atoms, but in some cases ranging from about 11 to about 16 or so carbon atoms. Propylene can be 10 dimerized to hexenes, and the hexenes can be converted to aldehydes by an oxo reaction as described herein, and the resulting heptaldehydes can be reacted in an aldol reaction to produce aldol products which can be hydrogenated to C₁₄ alcohols. Using oxo product 15 in which the content of aldehyde without 2-substitution may range from 60 up to near 80% or so, aldol conversions may approach 75 to 90%, even though participation of the 2-substituted aldehydes is limited, so that it is involved in cross-aldol 20 producing no more than 20% of the product.

The aldol reaction is carried out for the most part utilizing the usual aldol catalysts and temperature conditions, using elevated temperatures upwards of 60°C., particularly temperatures of about 25 90°C. to 130°C., or possibly up to 150°C. or higher if desired, the conditions also being those disclosed for n-pentanal in S.N. 272,587. The reaction is operable over broad pressure ranges including pressures less than atmospheric as well as elevated pressures, but 30 will usually be effected at slightly elevated pressures sufficient to maintain the reactants substantially in the liquid state. The reaction can also conveniently be conducted at reflux.

The aldol reaction can utilize strongly 35 alkaline catalyst, such as sodium and potassium



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hydroxide, or sodium and potassium cyanide. The concentration of the aqueous alkali can be varied, but molar or similar concentrations of alkali metal hydroxides can be used, and concentrations selected 5 will generally be in the range of about 1 to 10% by weight. The amount of aqueous alkali to aldehyde reactant can also vary, for example from about 15% by volume aqueous alkali up to about 75% by volume aqueous alkali. The aldol reaction will be run for a 10 sufficient time to obtain the desired degree of conversion, which for batch reactions may be in the range of about 1 to about 3 hours, while in continuous reaction times of less than five minutes are achievable. The reaction is stopped by permitting the 15 reaction mixture to cool and separating the organic reaction phase from the aqueous alkali phase.

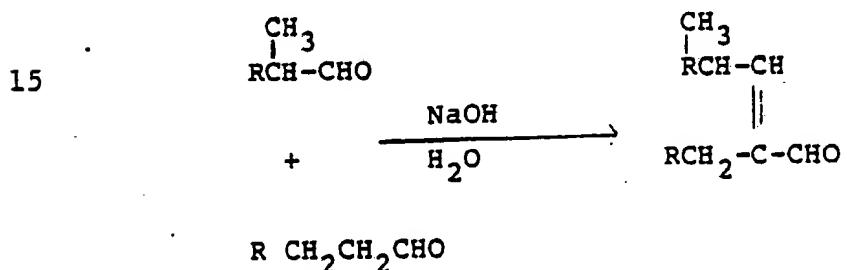
In one major respect it is difficult to conduct aldol reactions of heptanals in conventional manner, as in aqueous NaOH, because this procedure 20 requires appreciable solubility of the organic aldehyde in the aqueous phase, or vice versa. The mutual solubility is so low with C₇ aldehydes that little reaction occurs in reasonable time periods. However, it has been found that use of a co-solvent 25 overcomes this problem and results in suitable reaction rates. In principle, any solvent with miscibility with both the aldehyde and the aqueous base, or at least some solubility with respect to each, will act to increase the rate of the reaction. 30 It is also desirable that the solvent be relatively inert under the reaction conditions so as not to cause interfering reactions or to be readily degraded excessively by the hot basic medium. In general, polar solvents will tend to have the requisite 35 solubility characteristics, and hydroxy alkanes, for



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example, alkane diols, having the appropriate solubility characteristics can be used. Hexanediol is very suitable. Methanol is also a suitable cosolvent.

In the aldol reactions involved in the present invention it is necessary to utilize a high proportion of aldehydes without 2-branching, also referred to herein as aldolable aldehydes, in order to achieve good yields of aldol. The 2-substituted aldehydes do not undergo self-condensation aldol reactions with any facility under the usual basic aldol conditions, although they will serve as acceptor molecules to some extent in cross-aldol reactions, as illustrated with the following heptanal isomers:



in which R represents a butyl group. However, the 2-substituted aldehydes react more slowly than other aldehydes, and therefore would constitute an undesirably high residue of unreacted component if provided to the reaction mixture in high proportion. In the present process with 75 to 80% or so of aldehydes without 2-branching, the aldol reaction can be conducted to include some cross-aldol of 2-substituted aldehydes so as to have yields of 85% or more based on starting aldehyde. In general, the C₁₄



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aldehydes resulting from such cross-aldol τ actions, and the alcohols produced therefrom, have properties comparable to those produced from aldol reactions of aldehydes without 2-substitution, and are suitably 5 present in alcohol mixtures for detergent preparations as described herein. The conversion of aldehydes without 2-branching in such aldol reactions can be in the range of 95% or better while possibly only about 1/4 to 1/3 of the 2-substituted aldehydes are 10 converted to aldol product.

The hydrogenation of the enals from the aldol reaction can be conducted under the usual catalytic hydrogenation conditions for reducing olefinic bonds and aldehyde groups. The 15 carbon-to-carbon bond reduces more rapidly and at a lower temperature than the aldehyde group, e.g. at about 90°C., with cobalt on Kieselguhr catalyst at elevated hydrogen pressure. The hydrogenation will generally be carried out at 100-20000 psi, or greater 20 hydrogen pressures and temperatures of 100° to 200°C. or higher, although any temperatures which are effective with a particular catalyst can be used. The stated conditions will be effective for reducing both the carbon-to-carbon bond and the aldehyde group to 25 obtain saturated alcohol. Various other hydrogenation catalysts can be used including platinum and platinum on carbon catalysts, copper chromite, activated nickel, etc., and individual catalysts can be utilized in conjunction with other catalysts.

30 The present invention can include an oxo reaction, followed by an aldol reaction, and then a hydrogenation to convert enals to alcohols. For large scale operations, the oxo reaction will be conducted with usual provisions for separating gaseous reactants 35 and products, and catalysts, from the aldehyde



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products, with recycle as appropriate. The aldehyde product mixture will then be subjected to an aldol reaction, followed by decantation and water washing or other simple procedures to separate the organic 5 product-containing phase from the aqueous phase. The product phase is then hydrogenated, converting both unreacted heptanals and C₁₄ enals to the corresponding alcohols when the product phase is from a heptanal reaction. The hydrogenation is followed by a 10 distillation to remove light ends, followed by a distillation to remove C₇ alcohols. Both the C₇ and C₁₄ alcohols can then be treated in further hydrogenation polishing operations to improve the alcohol quality by insuring complete hydrogenation.

15 The separation of the C₇ from C₁₄ alcohols is readily effected by distillation in equipment constructed of inexpensive alloys such as carbon steel. Separation at this stage is simple, compared to the difficult separation which would be required to 20 separate the seven carbon aldehyde isomers prior to the aldol reaction.

As an alternate to the above procedure, it is possible to separate the unreacted seven-carbon aldehydes by distillation from the 14 carbon enals 25 prior to hydrogenation. For convenience of separation, distillation of the alcohols is generally preferred. However, if the aldehydes are desired for some purpose, separation is appropriate, and this has the advantage of avoiding unnecessary hydrogen use. 30 The aldehyde separation can, of course, be effected with other aldehydes, e.g. to separate five-carbon aldehydes from 10-carbon enals.

In the present process the oxo product is used in the aldol reaction without any need for 35 separation of some of the components. This contrasts



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with the usual commercial procedure, for example, for preparing 2-ethylhexanol as a plasticizer alcohol, wherein it is the practice to remove isobutanal before conducting an aldol reaction with n-butanal. The 5 separation is effected by distillation and, since isomeric aldehydes have similar boiling points, separation on a commercial scale involves high capital cost equipment with consequent expense, and a substantial energy cost. There is a definite 10 advantage in avoiding such a distillation step in the present process.

The C₇ branched aldehydes which do not react in the aldol reaction can be separated from the reaction mixture for various purposes, or hydrogenated 15 with the mixture and utilized as a C₇ branched alcohol. If desired, the branched C₇ alcohol can be dehydrated, and then subjected to an oxo reaction to produce an aldehyde with an additional carbon atom. Thus 2-methyl-hexanol can be converted to 20 2-methylhexene-1, which can be recycled to the oxo stage of the reaction process and hydroformylated to predominantly 3-methyl-heptanal. This 3-methylheptanal, not having any substituent in the 2-position, reacts at a good rate in the aldol 25 reaction. Other unreacted aldehydes, such as 2,4-dimethylpentanal and 2-ethyl-pentanal, can similarly be hydrogenated and dehydrated and recycled to be converted in the oxo stage to C₈ aldehydes with no substituent in the 2-position, which will take part 30 in the aldol reaction when recycled through that stage. This procedure to use the unreacted C₇ aldehyde results in greater conversion of the original reactants to the desired final product, rather than to a concomitant product such as C₇ alcohols. Use of 35 this recycle feature in the oxo-aldol process changes



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the components of the product from mixed hexenes to some extent, but the properties will be similar as the main difference in the additional components will be an additional methyl substituent as in 2(1-methylbuty)-5-methylnonanol, 2-(3-methylbuty)-5-methylnonanol and 2-pentyl-5-methylnonanol, for example.

5 The Dimersol [®] dimerization process has been referred to in various publications, e.g. see "How First Dimersol is Working" by Benedek et al, 10 Hydrocarbon Processing, May 1980, page 143; also Chauvin et al, "The IFP Dimersol [®] Process for the Dimerization of C₃ and C₄ Olefinic Cuts", Advances in Petrochemical Technology, presented at American Institute of Chemical Engineers, April 13, 1976, 15 Kansas City, Missouri.

The combination of the Dimersol [®] dimerization process, oxo process, aldol and hydrogenation provides a very efficient route from propylene to detergent range alcohols. One of the 20 known routes to such alcohols relies upon oligomerization of ethylene to obtain higher molecular weight materials which are then subjected to an oxo reaction. The presently proposed route is in many respects more efficient and economical than those 25 involving ethylene oligomerization, as propylene costs less than ethylene, and the reactions involved using dimerization, oxo and aldol are more straight forward than an oligomerization which can produce a broad mixture of products and require extensive equipment 30 and procedures to direct it to suitable product. As discussed hereinabove, the mixture of isomers obtained from a dimerization can be carried through the oxo, aldol and hydrogenation reactions to obtain high overall conversions and yields, despite the presence 35 of extensive branching in the materials. It is



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fortunate to find that a high proportion of the materials are capable of taking part sequentially in all of the required reactions, and in particular that the aldehyde failing to react to a significant degree 5 in the aldol reaction, because of 2-substitution, is at a comparatively low level.

In the oxo stage of the present process it will be noted that cobalt catalyst is employed with the hexenes in order to promote migration of the 10 olefinic bond and high selectivity to desired aldehyde isomers, such catalysts being for example $\text{Co}_2(\text{CO})_8$ which may be equivalent to $\text{HCo}(\text{CO})_4$ under reaction conditions.

The processes of the present invention can 15 utilize mixtures with olefins of different carbon numbers, along with isomeric mixtures of olefins of particular carbon number. The oxo product of such mixtures can be subjected to aldol reactions as taught herein to give aldol product. Such procedures can, 20 for example, use butenes in conjunction with Dimate^R hexenes. Also, Dimate^R hexenes may at times have various amounts of C_5 , C_7 and C_9 olefins, and still be usefully employed in the present invention. Such processes will still have the advantage of production 25 of high amounts of aldolable aldehydes in the oxo process, and result in aldol product with carbon numbers in the detergent range and with branched structure suitable for detergent use. Also, in the Dimersol^R process, butene can also be dimerized or 30 codimerized with propylene, and this provides additional ways to modify the present process by variation of the olefin feed stock, while still producing aldol product in the detergent range. In the procedures which use mixed olefins, it is also 35 possible to conduct the oxo reaction with olefins of



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different carbon number, and to combine the resulting oxo products for an aldol reaction. Another olefin of particular interest for use in the present process is 4-methyl-1-pentene, which can be produced by Na catalyst 5 dimerization of propylene, as reported in Industrial Organic Chemistry, cited hereinabove. The reaction is conducted at approximately 150°C and 40 bar, using Na/K₂CO₃. The 4-methyl-1-pentene in an oxo process will give very high aldolable product as shown by the 10 results for 4-methyl-pentene-2 in Example 4, and is therefore well suited for use in the present process to prepare C₁₄ enals and detergent derivatives thereof.

The C₁₄ or other detergent range alcohols 15 produced by the present process can be readily converted to detergents by known procedures. Thus non-ionic detergents are prepared by reaction with ethylene oxide to have a desired number of ethoxyl groups, e.g. 6 to 10 or 12 or so. These, or other 20 ethoxylated alcohols, possibly with 2 to 3 ethoxyl groups can be reacted to form an alcohol ether sulfate, having a sulfate anionic end group with a sodium or other cation. The alcohols can also be reacted to prepared sulfate derivatives. The 25 detergents thus prepared will have the requisite hydrophobic groups for detergent properties. Moreover, the structures are such as to provide biodegradability, in that the structures are acyclic alkyl groups which are essentially free of any 30 quaternary carbon groups. There is some branching on adjacent carbon atoms, but that and the common 2-branching characteristic of aldol product, with or without various additional methyl or other lower alkyl branches in non-adjacent positions, do not have any 35 important effect on the biodegradable nature of the



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compounds. An alcohol ether sulfate prepared from 2-pentylnonanol has been described as biodegradable by Crawland et al, Surfactant Congress No. 4, Vol. 1, page 93 (1967). Also Kravetz et al, Proceedings of the American Oil Chemists' Society, 69th annual meeting, May, 1978, St. Louis, MO., concluded that variation of branching from 45% to 75% linear had no appreciable effect on biodegradation rates of primary alcohol ethoxylates, and make reference to 58% branching giving biodegradation at rates not appreciably different from zero branching. The branching involved in both of these prior studies was a single branch, and this can be contrasted with the materials described herein which contain at least two branches in almost all cases.

A quantity of Dimate® hexenes from a refinery stream was distilled to have a C₆ cut, approximately 73% of the total material. Analysis is given in Table 1. The Dimate® hexenes had been produced by dimerization of propylene over a catalyst by the Dimersol® process.



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TABLE 1
Dimersol ® Hexene Distribution
 % (100% basis)

	2,3-dimethyl-2-butene	4.5
	2-methyl-2-pentene	35.6
5	trans-4-methyl-2-pentene	18.4
	cis-4-methyl-2-pentene	3.7
	2-methyl-1-pentene	5.1
	2,3-dimethyl-1-butene +	
	4-methyl-pentene-1	1.7
10	trans-2-hexene	17.8
	trans-3-hexene	6.3
	cis-3 + cis-2 hexene	6.8
	1-hexene	0.1
		31.0% linear

15 The distillation serves to remove some C₅, C₇ and C₉ hydrocarbons resulting from oligomerization involving some ethylene present in the original olefin feed, or trimerizations. It also removes C₆ chlorides, along with the C₉ hydrocarbons; these 20 chlorides, resulting from the dimerization catalyst, could contaminate the oxo catalyst if not removed.

25 It is fortunate that the hexenes mixture is amenable to reaction at a good rate in the oxo reaction. It is sufficiently reactive to permit use of moderate conditions and equipment therefor, with suitable reaction rates and times. This is in contrast to an octenes Dimate ® mixture which is characterized by more relatively unreactive dibranched olefin isomers and much slower reaction rates. Such 30 material requires more severe reaction conditions in more expensive equipment, and additional reactor capacity.



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Since the oxo product is to be reacted in an aldol reaction, it is to be conducted under conditions which favor production of aldehydes as contrasted with alcohol or other products.

5

EXAMPLE 1

Hydroformylation of a Dimate hexenes mixture, of composition reported in Table 1, was carried out in an autoclave with agitation. The autoclave was charged with 0.52 g dicobalt 10 octacarbonyl, 74.06 Dimate hexenes and 3000 psi. gauge (20,786 KPa) of 1:1 CO and H₂. The autoclave was heated to 130°C. and held for four hours. Liquid samples were taken every hour. The autoclave was cooled rapidly and the product removed under nitrogen. 15 Analysis indicated 92% conversion of the olefins. The product was analyzed chromatographically, with results as reported in Table 2 (along with other examples). It will be noted that 77.9% of the aldehyde product was unbranched at the 2-position and therefore aldolable. 20 This result was obtained, even though the hexene reactants were more than 90% composed of internal olefins, including some linear hexenes which give product little more than 50% aldolable. A similar run employing a 110°C. temperature with cobalt catalyst 25 for 23 hours is also reported.

EXAMPLE 2

The autoclave of Example 1 was charged with 0.49 g. of dicobalt octacarbonyl, 69.84 g. 2-methylpentene-2 and 3000 psi gauge (20,786 KPa) of 30 CO and H₂ in 1 to 1 ratio. The autoclave was heated to 130°C. and held at this temperature for three hours. The autoclave was cooled and the product removed under a blanket of argon, and analyzed. Conversion of the olefin was 83%. The procedure was 35 repeated substantially, but employing a temperature of



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116°C. for a 48% conversion. Results of the analysis are reported in Table 2. It is notable that better than 90% of aldehydes unbranched at the 2-position, i.e. aldolable aldehydes, were obtained.

5 Approximately the same results were shown from samples taken during the reactions, with the 116° reaction at 1 hour giving 17% conversion and 35.4% 5-methylhexanal and 55.5% 3-methylhexanal; and the 130° reaction temperature 45% conversion at 1 hour with 32.5% 10 5-methylhexanal and 57.6% 3-methylhexanal.

EXAMPLE 3

Hydroformylation was carried out on 2-methyl-1-pentene with cobalt catalyst in accord with the procedure of Example 2 at 116°C. for a three-hour 15 period. A 41% conversion was obtained, with 95% of aldolable aldehyde unbranched at the 2-position being obtained. Results are reported in Table 2.

EXAMPLE 4

Hydroformylation was effected with 20 4-methyl-2-pentene in accord with the Example 2 procedure, employing a 130°C. temperature. Results are reported in Table 2.



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Table 2
Oxo Product Distribution

Hexene	Temp. (°C)	Run Time (hours)	%									
			I	II	III	IV	V	VI	VII	VIII	IX	
0-Mixture	110	23	15.3	19.4	0.7	39.1	6.4	3.1	1.4	7.5	1.0	76.9
p-Mixture	130	6	16.9	25.0	0.5	36.2	4.3	1.0	1.1	7.4	0.8	77.9
" "	3	15.6	23.5	0.0	37.9	4.5	0.7	1.1	7.5	0.7	77.5	
" "	2	16.0	25.5	9.6	33.0	4.9	0.6	1.0	7.2	0.6	76.9	
" "	1	10.8	26.5	10.7	31.8	5.4	trace	1.4	6.8	0.5	75.1	
2-methyl-2-pentene	116	3	—	34.3	—	56.6	—	—	1.3	6.9	0.9	90.9
2-methyl-1-2-pentene	130	3	—	33.6	—	56.6	—	—	1.4	7.2	1.2	90.2
2-methyl-1-pentene	116	3	—	10.6	—	86.4	—	—	0.4	2.5	2.1	95.0
3-methyl-1-2-pentene	130	3	—	65.9	—	17.3	trace	1.6	15.3	trace	0.3	2.2
IV	2	—	65.1	—	16.6	—	—	1.6	16.8	trace	0.1	77.5
V	1	—	67.6	—	14.2	—	—	1.4	16.0	trace	0.1	81.8

VI = 3,4-dimethylpentanol

VII = 2,3-ethyl-3-methylbutanal

VIII = 2,4-dimethylpentanol

IX = 2,2-dimethylpentanal

I = heptanol

II = 5-methylhexanal

III = 2-methylhexane

IV = 3-methylhexanal

V = 2-ethylpentane



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EXAMPLE 5

Hydroformylation was effected with 2,3-dimethyl-2-butene at 130°C, employing the procedure of Example 2. At 1, 2 and 3 hours 5 conversions were respectively 18, 43 and 71%, and in each case the aldehyde product was analyzed as 100% 3,4-dimethylpentanal, an aldolable aldehyde.

EXAMPLE 6

This example was a simulation of two oxo 10 reactors operating at different temperatures. The autoclave was charged with 0.44 g. dicobalt octacarbonyl, and 73.94 g. Dimate hexenes. The autoclave was sealed, pressure checked and run under 3000 psig(20,786 KPa) of 1/1 CO to H₂ at 130°C for 1 15 hour then heated to 140°C for an additional 4 hours. The autoclave and contents were rapidly cooled and the product removed under nitrogen. Analysis of the final product yielded that the olefin was 97% converted with 83% as the aldehydes, 11% as alcohols, and 3% high 20 boilers. The unreacted olefin consists of 2-methylpentene-2, 2,3-dimethylbutene-2 and trans-hexene-2. The aldehydes after normalization are 39% 3-methylhexanal, 20% 5-methylhexanal, 16% heptanal, 9% 2-methylhexanal, 6% 2,4-dimethylpentanal, 25 5% 2-ethylpentanal, 4% 3,4-dimethylpentanal, 1% 2-ethyl-3-methylbutanal, 1% 2,2-dimethylpentanal. The aldehydes were distilled from the catalyst, unreacted starting material and other products at 20 mmHg from 41 to 45°C.

30 The percentage of aldehydes unbranched at the 2-position produced was 79%.

The results in Example 6 indicate that the percentage of aldehydes unbranched at the 2-position from branched hexenes (69% of hexenes) was 88.7%.



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compared to only 53.3% of such aldehydes from the linear hexenes (31% of hexenes). Considering only the methylpentenes (62.8%, ignoring the small amount with 2,4-dimethylbutene in Table 1), the aldolable aldehyde 5 unbranched at the 2-position content of the aldehyde product was 86.6%).

EXAMPLE 7

An aldehyde mixture representative of that from the oxo reaction of Dimate® hexenes, as 10 described in Example 16, was reacted in an aldol reaction. An autoclave was charged with 50 ml of 0.8 M NaOH, 100 ml of 2,5-hexanediol under 20 psi gauge argon. Then 36.2 g., 50 ml, of the aldehydes were pressured into the autoclave with argon after the 15 autoclave had been heated to 100°C. and agitation set at 1500 rpm. The reaction was run for an additional hour and the system rapidly cooled. The product was removed and the upper and lower phases were separated. The upper phase contained 16.8% unreacted aldehydes, 20 2.4% heptanols, 72.9% tetradeconals and 4.1% 2,5-hexanediol. The conversion of the aldehydes approximated 79%.

EXAMPLE 8

Unsaturated aldehydes representative of 25 product from oxo and aldol reactions of Dimate® hexenes as produced in Example 7, were subjected to hydrogenation. An autoclave was charged with 130.95 g of 45% + 5% cobalt on Kieselguhr, 1336.1 grams of the aldol condensation product and hydrogen to 1500 psi 30 gauge (10,644 KPa). The autoclave was carefully heated to 160°C., with gas uptake starting at 100°C. The pressure and temperature were maintained for 4 1/2 hours. The catalyst was filtered off and washed with methanol to remove any residual alcohols. The 35 material was then distilled using a 30 cm. vigreux



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column at 5 mm Hg. The C₁₄ alcohols were collected from 119 to 122°C. From analysis of the aldehydes prior to the aldol reaction, and analysis of derivatives of some of the alcohol components, 5 together with gas chromatographic and other identification as included and described hereinbelow, it was concluded that the product includes some 28 alcohols in amounts and as named and illustrated by skeletal structure in Table 3 where the 10 hydroxyl-bearing group is designated by an asterisk.



Table 3

tetradecanols

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<chem>CCCCCCCCC</chem>	2-(1-methylbutyl)-7-methyloctanol	2.4
<chem>CCCCCCCCC</chem>	2-pentyl-4-methyloctanol	2.4
<chem>CCCCCCCCC</chem>	2-pentyl-5,6-dimethylheptanol	2.3
<chem>CCCCCCCCC</chem>	2-(3-methylbutyl)-4-methyloctanol	2.2
<chem>CCCCCCCCC</chem>	2-(1-methylbutyl)-nonanol	1.5
<chem>CCCCCCCCC</chem>	2-(1-methylbutyl)-nonanol	1.5
<chem>CCCCCCCCC</chem>	2-(1-methylbutyl)-4-ethylheptanol	0.7
<chem>CCCCCCCCC</chem>	2-pentyl-4-ethylheptanol	0.6
<chem>CCCCCCCCC</chem>	2-(1,2-dimethylpropyl)-nonanol	0.5



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<chem>CCCCCCCCC</chem>	2-(1-methylbutyl)-4-methyloctanol	0.4
<chem>CCCCCCCCC</chem>	2,(1,2-dimethylpropyl)-5,6-dimethyl-heptanol	0.4
<chem>CCCCCCCC</chem>	2-(3-methylbutyl)-4-ethylheptanol	0.2
<chem>CCCCCCCC</chem>	2-(1,2-dimethylpropyl)-7-methyl-octanol	UNK
<chem>CCCCCCCC</chem>	2-(1,2-dimethylpropyl)-5-methyl-octanol	UNK
<chem>CCCCCCCC</chem>	2-(1,2-dimethylpropyl)-4-methyl-octanol	UNK
<chem>CCCCCCCC</chem>	2-(1,2-dimethylpropyl)-2-ethyl-heptanol	UNK
<chem>CCCCCCCC</chem>	2-pentyl-4,6-dimethylheptanol	UNK
<chem>CCCCCCCC</chem>	2-(3-methylbutyl)-4,6-dimethyl-heptanol	UNK

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UNK

2-(1,2-dimethylpropyl)-4,6-
dimethylheptanol

C C C C C C C
C C C* C C C
C

UNK

2-(1-methylbutyl)-4,6-
dimethylheptanol

C C C C C C C
C C C* C C C
C



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5 Chromatographic retention times for particular alcohols, both as components of a mixture and from individual synthesis, are set forth in Table 4, together with the aldehyde pair involved in the synthesis.

10 Each of the individual alcohols in Tables 3 and 4, as well as the total mixture, can be converted to ethoxyl derivatives, sulfates, or ethoxyl sulfate derivatives. This is also the case with other alcohols described herein. Similarly, the individual enals and enal mixtures can be readily converted to saturated aldehydes, alcohols and acids.



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TABLE 4
TETRADECANOLS GC IDENTIFICATION

Name	Retention Time In Mixture	Aldol, Condensation Substrates Carbanion Acceptor	Retention Time From Individual Synthesis
2-(3-methylbutyl)-5-methyloctanol	31.99	5-methylhexanal	32.36
2-(1-methylbutyl)-5-methyloctanol	32.55 +	3-methylhexanal	32.43 +
2-pentyl-5-methyloctanol	32.74	3-methylhexanal	32.60
2-(3-methylbutyl)-7-methyloctanol	36.62	Heptanal	36.49
2-pentylheptanol	33.32	5-methylhexanal	33.27
2-(3-methylbutyl)-5,6-dimethylheptanol	43.60	Heptanal	43.44
2-(1-methylbutyl)-5,6-dimethylheptanol	31.27 31.27	5-methylhexanal 3-methylhexanal	31.24 31.28
2-pentyl-7-methyloctanol	37.86 or 38.18	Heptanal 5-methylhexanal	37.93 38.29 or
2-(3-methylbutyl)-nonanol	37.86 or 38.18	5-methylhexanal Heptanal	37.93 38.29 or
2-(1-methylbutyl)-7-methyloctanol	34.17 +	3-methylhexanal	34.31 +
	34.40		34.50



TABLE 4 (Continued)
TETRADECANOLS GC IDENTIFICATION

<u>Name</u>	<u>Retention Time In Minutes</u>	<u>Aldol Condensation Substrates</u>		<u>Retention Time From Individual Synthesis</u>
		<u>Carbonyl Acceptor</u>	<u>Carbonyl Substrate</u>	
2-pentyl-4-methyloctanol	34.17	Heptanal	2-methylhexanal	34.15
	+		+	+
	34.40			34.41
2-pentyl-5,6-dimethylheptanol	35.79	Heptanal	3,4-dimethylpentanal	35.82
	29.85			29.84
2-(3-methylbutyl)-4-methyloctanol	+	5-methylhexanal	2-methylhexanal	+
	30.11			30.09
	38.98			38.89
2-(1-methylbutyl)-nonanol	+	3-methylhexanal	Heptanal	+
	39.13			39.04
	26.67			26.67
2-(1-methylbutyl)-4-ethylheptanol	+	3-methylhexanal	2-ethylpentanal	+
	26.95			27.05
2-pentyl-4-ethylheptanol	33.05	Heptanal	2-ethylpentanal	32.97
2-(1,2-dimethylpropyl)-nonanol	30.11	3,4-dimethylpentanal	Heptanal	30.58
	30.40			30.38
2-(1-methylbutyl)-4-methyloctanol	+	3-methylhexanal	2-methylhexanal	+
	30.84			30.83

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TABLE 4 (Continued)
TETRADECANOLS GC IDENTIFICATION

Name	Retention Time In Minutes	Aldol Condensation Substrates	Retention Time From Individual Synthesis
		Carbanion	Acceptor
2-(1,2-dimethylpropyl)-5,6-dimethylheptanol	27.82 +	3,4-dimethylpentanal	27.79 +
	28.86	+	28.73
2-(3-methylbutyl)-4-ethylheptanol	29.13 28.86 +	5-methylhexanal	29.01 28.87 +
	28.97	2-ethylpentanal	28.89
2-(1,2-dimethylpropyl)-7-methyloctanol		3,4-dimethylpentanal	5-methylhexanal
2-(1,2-dimethylpropyl)-5-methyloctanol		3,4-dimethylpentanal	3-methylhexanal
2-(1,2-dimethylpropyl)-4-methyloctanol		3,4-dimethylpentanal	2-methylhexanal
2-(1,2-dimethylpropyl)-2-ethylheptanol		3,4-dimethylpentanal	2-ethylpentanal
2-pentyl-4,6-dimethylheptanol		2-ethylpentanal	2,4-dimethylpentanal
2-(3-methylbutyl)-4,6-dimethylheptanol		5-methylhexanal	2,4-dimethylpentanal
2-(1,2-dimethylpropyl)-4,6-dimethylheptanol		3,4-dimethylpentanal	2,4-dimethylpentanal
2-(1-methylbutyl)-4,6-dimethylheptanol		3-methylhexanal	2,4-dimethylhexanal

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Carbanion and acceptor columns present the two aldehydes that were used in aldos to prepare the respective alcohol.



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The identification and reported percentage composition of the product alcohols were accomplished by utilizing gas-liquid chromatography (G.C.) coupled with synthesis of the isomers. The individual 5 alcohols were synthesized by aldol condensation of pure heptanal aldehyde isomers followed by hydrogenation to the alcohol. The heptanal pairs can be reacted together with the procedure of Example 7 above. The elution time of the individual derivatized 10 isomers were then matched with peak-elution times in the product mixture to identify the compounds in the mixture. Verification of the structures was accomplished by several methods. Gas chromatography combined with mass spectrometry (G.C. - MS) was used 15 to show that the mixture contained only tetradecanol isomers. The logic by which we synthesized the various alcohol isomers, discussed above, was confirmed in three cases by ^{13}C nuclear magnetic resonance (nmr) and also in one case by a single 20 crystal X-ray structural determination of a solid derivative. Thus, the structures of 2-pentylnonanol, 2-(1-methybutyl-5-methyl-octanol and 2-(3-methylbutyl) 25 7-methyloctanol were unambiguously identified using two dimensional double quantum coherence ^{13}C nmr techniques (G. Bodenhouse, "Progress in NMR Spectroscopy," 14, 137 (1981)). In addition, the -napthyl sulfonate ester of 2-(3-methylbutyl-7-methyl 30 octanol was prepared (m. pt 47°) and a single crystal suitable for X-ray crystallographic investigation was grown. A single crystal study was performed using copper K radiation with a Syntex P2₁ diffractometer. The space group was found to be P2₁/N. Cell constants were = 8.186(1), b = 29.958 (7), c = 10.316(2) Å; a = 107.49(2)°; z = 4; M = 404.62. The structure was



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solved by direct methods and refined to a final R_{f} value of 0.091 using 2586 observed reflections. The structure is shown in the accompanying Figure 1. The bond lengths and bond angles are given in Table 5.

This X-ray determination unequivocably
5 confirms the backbone structure of the alcohol as
2-(3-methylbutyl)7-methyl octanol.

TABLE 5

Bond Lengths and Bond Angles in the -Naphthyl
Sulfonate Ester of 2- (3-methylbutyl-7-methyl Octanol
10 (Numbering system used here is shown in Fig 1)

Bond Lengths

	S1	-	O2	1.565 Å
15	S1	-	O11	1.427
	S1	-	O12	1.427
	O2	-	C3	1.493
	C3	-	C4	1.543
	C4	-	C5	1.526
20	C4	-	C23	1.538
	C5	-	C6	1.534
	C6	-	C7	1.542
	C7	-	C8	1.544
	C8	-	C9	1.573
30	C9	-	C10	1.574
	S1	-	C14	1.747
	C13	-	C14	1.392
	C13	-	C18	1.399
35	C14	-	C15	1.421
	C15	-	C16	1.347
	C16	-	C17	1.416
	C17	-	C22	1.417
	C18	-	C19	1.442



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	C20	-	C21	1.449
	C21	-	C22	1.372
	C23	-	C24	1.545
	C24	-	C25	1.564
5	C25	-	C26	1.540
	C25	-	C27	1.393

Bond Angles

10	02	-	S1	-	011	104.1°
	02	-	S1	-	012	109.8
	02	-	S1	-	C14	103.3
	011	-	S1	-	012	119.9
	011	-	S1	-	C14	108.9
15	012	-	S1	-	C14	109.5
	S1	-	02	-	C3	117.9
	02	-	C3	-	C4	106.0
	C3	-	C4	-	C5	111.4
	C3	-	C4	-	C23	113.2
20	C5	-	C4	-	C23	109.4
	C4	-	C5	-	C6	115.8
	C5	-	C6	-	C7	110.1
	C6	-	C7	-	C8	110.6
	C7	-	C8	-	C9	111.7
25	C8	-	C9	-	C10	112.5
	C8	-	C9	-	C28	106.8
	C10	-	C9	-	C28	112.4
	C14	-	C13	-	C18	118.6
30	S1	-	C14	-	C13	119.9
	S1	-	C14	-	C15	118.6
	C13	-	C14	-	C15	121.6
	C14	-	C15	-	C16	119.6
	C15	-	C16	-	C17	121.5



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	C16	-	C17	-	C18		118.4
	C16	-	C17	-	C22		122.5
	C18	-	C17	-	C22		119.1
	C13	-	C18	-	C17		120.3
	C13	-	C18	-	C19		120.0
5	C17	-	C18	-	C19		119.6
	C18	-	C19	-	C20		119.8
	C19	-	C20	-	C21		120.2
	C20	-	C21	-	C22		120.6
	C17	-	C22	-	C21		120.6
10							
	C4	-	C23	-	C24		113.6
	C23	-	C24	-	C25		112.1
	C24	-	C25	-	C26		108.2
	C24	-	C25	-	C27		119.2
15	C26	-	C25	-	C27		114.3

In Fig. 1, the β -naphthylsulfonate ester of 2-(3-methylbutyl)-7-methyl octanol is shown. the sulfur and oxygen atoms are identified by as S and O respectively, while all other atoms are carbon. The portion of the structure below the oxygen atom 2 is the carbon structure of the alcohol, while that above 2 is the naphthylsulfonate portion of the derivative.

The gas chromatography for the retention procedures discussed was done with acetate ester derivatives. The acetate esters were prepared by utilizing 0.1 cc of alcohol sample, 0.3 cc acetic anhydride, 1 drop of pyridine, and heating in a sealed vial for 10 minutes at 100°C. The product was injected directly into a gas chromatogram, which was a Varian 3700 with splitter for glass capillary columns, Flame ionization detector and Hewlett-Packard 3352



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computer. Details of the column and procedure were:

SP-2100 Glass capillary column 60 meter length x 0.25 mm internal diameter, 27.1 cm/sec helium;

5 Injection/splitter 280°C.
Detector 280°C.
Column oven Progress from 120°C to
 150°C at 2°C/minute, and held
 at 150°C

10 Detector Sensitivity Range 10⁻¹¹ amps/millimole
 Injection Size 0.06 ul/split 60:1

15 A typical chromatograph of the acetate derivative of the mixture of alcohols of Table 3, obtained by the foregoing procedure, is shown in Figure 2 in which retention time is measured on the horizontal axis in minutes. The chromatograph was made at a chart speed of 5 mm/minute and covers from time zero to over 50 minutes. The designations of several peaks by combinations of two letters indicate the aldehydes combined in the aldol stage of the product formation, with the letters symbolizing:

25 A = n-heptanal
 B = 2-methylhexanal
 C = 3-methylhexanal
 D = 4-methylhexanal
 E = 2-ethylpentanal
 F = 2,3-dimethylpentanal

EXAMPLE 9

This example demonstrates an aldol condensation of



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heptanal in glassware under conventional aldol conditions. The reaction flask was charged with 116 ml of 0.8 M NaOH and the heptanal, 371.03 g., placed into the addition funnel. The system was kept under a nitrogen blanket and the agitation was set at 500 rpm.

5 The reaction flask was heated to reflux, and the heptanal added at a constant rate such that the addition was completed after 230 minutes. A sample was removed to compare with conventional aldol results and only 33.5% of the upper phase of the sample was

10 C_{14} unsaturated aldehyde and 61% was unreacted heptanal. The system was held at reflux for an additional 10 hours after which the system was cooled. The lower phase was removed, 135.74 g, and the upper phase, 336.06 g., was found to contain 66.5%

15 2-pentynon-2-enal, 26.4% unreacted heptanal and 2.5% heptanol.

EXAMPLE 10

This example demonstrates the efficacy of using a co-solvent with the water to enhance the aldol reaction of heptanal. The reaction flask was charged with 31 ml of 0.9 M NaOH and 100 ml methanol. The addition funnel was charged with 115 g. of heptanal under a nitrogen blanket. The aqueous-methanol mixture was heated to reflux with agitation of 450 rpm. The heptanal was added over a 40 minute period and refluxed for an additional 90 minutes. The system was cooled and the lower phase removed. The upper phase contains 10.1% methanol, 84.0% 2-pentynon-2-enal, 2.0% unreacted heptanal and 1.2% heptanol. This represents 94% conversion with 98% selectivity. The tetradecenals were hydrogenated as in Example 8 and the resulting saturated alcohols were treated as in example 8 including filtration and distillation.



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The use of methanol in this procedure was effective as a cosolvent, as compared to Example 9 where a cosolvent was not used.

EXAMPLE 11

5 Dimerization of propylene over transition metal catalysts produces a mixture of hexenes.

The linear hexenes in the mixture can be separated and such linear hexenes are for the most part 2-hexenes. The following procedure illustrates the 10 reaction of linear 2-hexenes in an oxo reaction, followed by an aldol reaction of the product. A sample of refinery 2-hexenes was passed through basic alumina particles for removal of oxides and 802 grams of the hexenes was placed in a 1 gallon stainless 15 steel autoclave with 4.75 grams $\text{Co}_2(\text{CO})_8$ catalyst. The autoclave was pressured to 2400 psi gauge with 1:1 carbon monoxide and hydrogen, and heated to 110°C . The temperature was kept at about 110°C for 1 hour and then rose to about 130° as the procedure was continued 20 for about 5 hours. A 916.5 gram product was obtained, with conversion about 94% with about 78% selectivity to C_7 compounds, and 71.5% to C_7 aldehydes. Chromatography indicated the aldehydes were in ratio 25 of about 29.4 n-heptanal to 16.7 2-methylhexanal to 8.5 2-ethylpentanal. A 907 gram amount of the product was distilled, with a final pot temperature of 120°C . and vacuum of 3 mm Hg. to obtain a 608 gram distillation fraction and 278 gram residue. Chromatography indicated the fraction included C_7 30 aldehydes in ratio 33.4 n-heptanal to 26.8 2-methylhexanal to 17.2 2-ethylpentanal, and minor amounts of other components. Evidently there was more loss of the normal aldehyde than the branched ones in the distillation.

35 It is feasible to achieve a higher percentage of



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n-aldehyde than present in the above distillation fraction, such as 60% or better, and therefore n-heptanal was added to the above fraction to have a more typical aldehyde for aldol reaction about 600 grams of the above fraction being used with 500 grams n-heptanal. A 300 ml amount of 0.8 molar sodium hydroxide was placed in a reaction flask with 955 ml methanol, and the aldehydes were placed in an addition funnel. The reaction medium was heated to about 71°C., and addition was slowly started and completed in about 13 hours. Chromatography indicated about 50% completion of the reaction, with C₁₄ aldehydes in ratio of about 23.4% 2-pentynon-2-enal to 8.72% 2-pentyl-4-methyloct-2-enal to 1.4% 2-pentyl-4-ethylhept-2-enal. Several C₇ aldehydes were also present in the ratio of 22.0 heptanal to 8.5 2-methylhexanal to 5.3 2-ethylpentanal.

The aldol condensation product was hydrogenated over a cobalt on Kieselguhr catalyst, using 131 grams catalyst with 1336 grams of the condensation product. The materials were maintained at about 160°C. and 1500 psi gauge (10,645 KPa) of hydrogen for about two hours when reaction appeared complete. Reaction conditions were maintained for an additional 4.5 hours. Analysis indicated about 99% completion of the hydrogenation. The product contained 2-pentyl-nonanol in about 18.4 to 7.2 ratio to a mixture of 2-pentyl-5-methyl-octanol and 2-pentyl-4-ethyl-heptanol and large amounts of C₇ alcohols from the unreacted aldehyde, being heptanol in a 27.5 to 16.3 ratio to a mixture of 2-methylhexanol and 2-ethylpentanol. The product was fractionated by distillation, with a 280 gram fraction being obtained at 110-115°C. at 2 mm Hg from 1180 grams of hydrogenation product. The fraction was in large predominance composed of C₁₄ alcohols.



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EXAMPLE 12

A mixture of hexenes produced by the Dimersol® dimerization process was utilized as olefin reactant. The crude hexene cut from the dimerization was used, and had the distribution of linear and branched hexenes typical of such material. A 1029 gram amount of the hexenes was used in a 1 gallon autoclave with 6.04 grams catalyst, $\text{Co}_2(\text{CO})_8$, 0.02 weight %. Peroxides had been removed from the hexenes by treatment on a basic alumina column. The autoclave was taken to reaction conditions with 1:1 CO/H_2 and maintained at 110°C . and 2600 psi gauge (18028 KPa) for 9 hours, with 80% of theoretical gas uptake, and then continued overnight. Chromatography indicated high conversion to C_7 aldehydes, with minor amounts of residual hexenes. A 1360 gram amount of the product was subjected to distillation, with a 797 gram fraction being obtained at pot temperatures of 60 to 97°C . as the vacuum dropped from 90 mm Hg to 5 mm Hg. Chromatography indicated a high portion of C_7 aldehydes with a very small amount of C_6 olefins.

A 792 gram amount of the above aldehyde fraction was utilized in an aldol reaction, adding the aldehyde material from an addition funnel to a reaction flask containing 564 grams methanol and 250.9 grams 0.8 molar sodium hydroxide. The addition took 6 hours, with stirring at about 500 rpm and temperature at $72-73^\circ\text{C}$. The reaction mixture was then refluxed for 1.5 hours. Analysis of a sample indicated only about 1 part aldol product to 3 parts aldehyde, on a mole basis. The reaction was continued at reflux overnight, giving 1 part aldol product to about 2.6 parts aldehyde reactant. During the reaction it was observed that the reaction mixture had a large upper phase and a smaller lower phase, indicating that



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methanol was not very effective in promoting miscibility and reaction, possibly because of the relatively long chain length of C₇ aldehydes.

5 Chromatography showed a fair amount of the C₁₄ aldol product, including 2-pentynonenal, and a large amount of unreacted C₇ aldehydes. (Results were better in Example 10 above in which a higher proportion of methanol was employed).

10 The above aldol product was subjected to further aldol reaction, after removing the methanol to employ different conditions. A 552 gram amount of the aldol condensate, 55.6 area percent C₇ aldehydes and 31.5 area percent C₁₄ enals, was placed in an addition funnel and added to a reaction flask containing 163 grams 0.8M NaOH and 389 grams 2,5-hexanediol.

15 Addition was complete after 45 minutes, with temperature maintained at 100°C. with agitation of the reaction mixture. The reaction mixture was then refluxed at 100°C for 1.75 hours. The reaction mixture separated into upper and lower phases of about equal weight. The conversion had been improved in that the ratio of C₁₄ enals to C₇ aldehydes in the product (upper phase) was now about 1.7 to 1.

20 A 515 gram amount of the product was subjected to hydrogenation, employing 51.65 grams cobalt on Kieselguhr catalyst and 160°C., about 1580 psi gauge (10,995 KPa) hydrogen. Approximately 549 grams of product was recovered. The conversion of C₁₄ saturated alcohols was about 90%, with about 10% found as unsaturated alcohols. The product was filtered to remove catalyst, and the filtrate was distilled. The process produced several C₁₄ alcohols in very substantial amounts, with a number of others in very small amounts. Several C₇ alcohols from unreacted aldehyde were also present in substantial amount.



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EXAMPLE 13

A freshly distilled sample of 2-pentene was hydroformylated in a 300 ml autoclave, employing 0.41 gram Co(CO)₈ catalyst with 65.84 grams pentene. A 1:1 mixture of CO/H₂ was used, with initial charge to 1500 psi gauge and heating to 120°C. and 3000 psi. with agitation at 1000 rpm. Gas-uptake was observed, as the pressure was increased to 3000 psi (20,684 KPa) After about 2 hours, an 83 gram product was obtained.

10 Chromatography showed a small residual amount of pentene and C₆ aldehydes in the ratio of 61.1 hexanal to 28.9 2-methylpentanal to 10.0 2-ethylbutanal. An aldehyde sample was provided to have aldehydes in the same ratio, using 189.1 grams hexanal, 89.9 grams 15 2-methyl-pentanal, and 31 grams 2-ethylbutanal, and placed in an addition funnel for addition to 110 ml of 0.8 M NaOH in a round bottom flask equipped with a mechanical stirrer. Heating was begun and addition was started after about 15 minutes and continued as reflux started around 91°C. Addition was completed in about 40 minutes. Stirring was continued for an hour, but without further heating, and a sample was taken.

20 Analysis indicated partial reaction. The reaction mixture was heated to 95°C. for an additional 1 1/2 hours. Upper and lower phases of the reaction mixture were separated, and the upper phase was analyzed. The analysis indicated better than 25% conversion to C₁₂ enal, nearly all being 2-butyloctanal, and large amounts of unreacted C₆ aldehydes, the major part of 25 which was branched aldehydes. The aldehyde mixture 30 can readily be hydrogenated to the corresponding alcohols.

EXAMPLE 14

An aldol procedure was carried out as in Example 35 13 except that the amount of water was increased ten



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fold. The same amount of NaOH was present, although now in much more dilute solution. Because of the large volume, less effective stirring was achieved. After a two hour reaction, analysis indicated substantial conversion to C₁₂ enals, although somewhat lower than in Example 12.

EXAMPLE 15

An aldol reaction was carried out as in Example 13, employing 6.1 to 2.9 to 1 ratio of hexanal to 2-methylpentanal to 2-ethylbutanal, the total amount being 310 grams. The branched aldehydes were placed in a flask with 110 ml of 0.8 M NaOH, and tetrabutylammonium chloride in an amount molecularly equivalent to the NaOH. The tetrabutyl-ammonium chloride serves as a phase transfer catalyst. The reaction flask was heated to reflux and addition was started. The addition continued for about 6 hours with reflux temperatures (pot) from 90-95°C. The mixture was cooled and separated into two phases. Analysis of the upper phase showed better than 75% conversion to C₁₂ enals, about half of which was 2-butyloctenal, and the remainder mainly a mixture of 2-butyl-4-ethyl-heptenal and 2-butyl-4-ethylhexenal. In the unreacted C₆ aldehydes present, the branched aldehydes were in greater amount the hexanal. The reaction can be directed to produce a higher percentage of product from the n-hexanal by adding the aldehydes together, rather than adding the n-aldehyde to the branched aldehydes in the reaction mixture as in the foregoing procedure. The phase transfer catalyst was effective in improving conversion in this procedure, but use of co-solvents, such as methanol or diols, may be more practical for large scale continuous operations. The use of hexanediol has been shown effective for aldol reaction of heptanals



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herein, and can similarly be used with hexanals.

It will be noted that the alcohols produced from both the pentenes and hexenes feedstocks are intended for use as detergent range alcohols. The 5 considerations herein as to reaction conditions and various parameters of the oxo and aldol reactions as described for the hexenes and resulting C₇ aldehydes also are in general applicable to the pentenes and resulting C₆ aldehydes. The C₁₂ alcohols produced 10 from the reactions starting with pentenes will have the hydrophobic groups such alcohols provide in detergents, and the groups will have a degree of branching similar to that of C₁₄ alcohols from hexenes. It is feasible to substantially avoid 15 presence of branches on adjacent carbon atoms. In one particular aspect the present invention is directed to a process of preparing alcohols from an oxo reaction with olefins selected from those having 5 to 6 carbon atoms, or mixtures thereof, to obtain aldehydes having 20 6 to 7 carbon atoms, comprising high amounts of aldehydes without 2-substitution and effecting aldol conversion with limited participation of 2-substituted aldehyde to obtain aldol product, which is then hydrogenated to C₁₂ or C₁₄ alcohols having properties 24 valuable for use in detergents.

EXAMPLE 16

An aldol reaction was conducted in a 1 liter round bottom flask equipped with stirrer, addition funnel, reflux condenser and adaptors for nitrogen flow. A 30 100 ml. amount of aqueous 1 molar potassium hydroxide solution was placed in the flask and heated to about 85°C. n-Pentanal and 2-methylbutanal were admixed in about 3:1 ratio, after each had been purified by distillation, and used for gradual addition to the 35 reaction flask with stirring. Over about one hour,



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about 300 ml was added containing 187.7 grams n-pentanal and 60.8 grams 2-methylbutanal. The reaction mixture was placed in a separatory funnel; and the lower aqueous phase (87 grams) was separated.

5 The organic layer was washed four times with water and amounted to 213.3 grams. Gas chromatographic analysis for the starting aldehydes indicated about a 3:1 ratio of 2-methylbutanal to n-pentanal, showing that the n-pentanal had been consumed at a much higher rate in the reaction. The product contained about 70.5% of 10 alkenal condensation product and 25.3% of the starting aldehydes. The product had 2-propyl-heptanal in about 9:1 ratio to 2-propyl-4-methyl-hexanal.

15 A 110.89 gram amount of the product was utilized for hydrogenation, employing 11 grams of cobalt on Kieselguhr catalyst with 4.4 ml H₂O as promoter, in a 300 ml stirred autoclave. The autoclave was pressured to 1000 psi with hydrogen and gradually heated, with hydrogen uptake starting at about 40°C. After one 20 hour, the pressure had fallen to 480 psi, and the autoclave was again pressured to 1000 psi. After two hours, with further addition of hydrogen, the pressure was 1510 psi and temperature 160°C. The run was continued for a total of sixteen hours. The measured 25 gas uptake was in very slight excess of theory for hydrogenation of both the olefin and aldehyde groups in the compounds present.

30 The product was filtered through a Celite filter mat to remove catalyst, and the mat was washed with n-hexane. The n-hexane was removed under vacuum, leaving 88 grams of product for distillation. Distillation was carried out at 10 mm Hg., with 18.1 gram being collected at 30-100°C., which gas chromatography indicated to be 66.4% 2-methylbutanol, 35 27.5% pentanol, and 4.1% 2-propylheptanal. An



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additional 38.9 grams was collected at 103.5-105°C., 11.3% 2-propyl-4-methylhexanol and 87.7% 2-propylheptanol. It can be seen that the above described procedure provides 2-propylheptanol with only very minor adulteration by the aldol alcohols product of branched aldehydes. Also the 2-propylheptanol from the 2-methylbutanol produced by hydrogenation of the 2-methylbutanal which did not undergo the aldol condensation.

10 Another sample of hexenes product from a Dimersol® dimerization refinery product was analyzed and found to have the following distribution:

Hexene Distribution

	% (100% Basis)
15	2,3-dimethyl-2-butene
	6.4
	2-methyl-2-pentene
	39.2
	trans-4-methyl-2-pentene
	15.9
	cis-4-methyl-2-pentene
	2.9
20	2-methyl-1-pentene
	5.0
	2,3-dimethyl-1-butene +
	4-methyl pentene-1
	1.7
	trans-2-hexene
	16.5
	trans-3-hexene
	5.8
25	cis 3 + cis 2-hexene
	5.6
	1-hexene
	1.0
	28.9%

The hexenes are suitable for conversion to detergent alcohol in accord with the present invention.

30 EXAMPLE 17

An aldehyde mixture representative of that from the oxo reaction of Dimate® hexenes, as described in Example 1, was reacted in an aldol condensation. A



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stirred autoclave was charged with 735 ml 0.8M NaOH, 1470 ml methanol and 20 psig argon (KPa) .The autoclave was heated to 100°C and 735 ml (551.0g) of the aldehydes were pressurized into the autoclave with 5 argon. The reaction was run for an additional hour and the system was rapidly cooled. The product was removed and the upper and lower phases were separated. The upper phase contained 14.4% unreacted aldehydes, 2.1% heptanols, 4.4% methanol, 10 1.6% high boilers and 77.5% tetradecenals. The tetradecenals product consists of some 28 isomers in amounts as named and illustrated in Table 5 where the aldehydic function is designated by an asterisk.

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TABLE 5
2-TETRAOCTENIIS FROM OXO AND ALDOL OF DIMATE HEXENYL

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3.4-BENZYL-2-CHLORO-1,1-DIMETHYL-2-GRIGNARD

2-METHYLHEXANAL	2-METHYLHEXANAL
2.2	5-METHYLHEXANAL

$ \begin{array}{c} \text{C} \\ \\ \text{C}-\text{C}-\text{C}-\text{C}-\text{C}-\text{C}-\text{C}-\text{C}-\text{C} \\ \quad \\ \text{C} \quad \text{C} \end{array} $	$(E+Z)-2-(1-\text{METHYLBUTYL})-2-\text{HEXENAL$	1.5 3-METHYLHEXANAL	MEPTANAL
------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------	--------------------------------------------------	--------------------------------------	-----------------

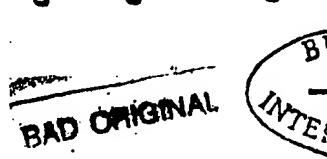
6.7	3-METHYLMETHANAL	(E+Z)-2-(1-METHYLBUTYL)-4-ETHYL-2-HEPTENAL	C
		C-C-C-C-C=C-C-C-C-C	

$$\begin{array}{c}
 \text{C} \\
 | \\
 \text{C}-\text{C}-\text{C}-\text{C}-\text{C}-\text{C}-\text{C}-\text{C} \\
 | \\
 \text{C} \\
 \text{C}=\text{C}-2-(1-\text{METHYLUTYL})-4-\text{METHYL}-2-\text{OCTENAL} \\
 \text{C} \\
 0.4 \quad 3-\text{METHYLBENZAL} \\
 \text{C} \\
 2-\text{METHYLHEXAL}
 \end{array}$$

2-ETHYLHEPTANAL	5-ETHYLHEPTANAL	0.2	(E+Z)-2-(3-METHYLBUTYL)-4-ETHYL-2-HEPTENAL	C-C-C-C-C-C-C-C-C-C
				1 1 1 1 1 1 1

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$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \text{C} \quad \text{C} \quad \text{C} \quad \text{C}=\text{C} \quad \text{C}-\text{C}-\text{C} \\ \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \end{array}$	(E+Z)-2-(1,2-DIMETHYLPROPYL)-7-HEXYL-2-OCTENAL	UNK	3,4-DIMETHYLPENTANAL	5-METHYLPENTANAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \text{C} \quad \text{C} \quad \text{C} \quad \text{C}=\text{C} \quad \text{C}-\text{C}-\text{C} \\ \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \end{array}$	(E+Z)-2-(1,2-DIMETHYLPROPYL)-5-METHYL-2-OCTENAL	UNK	3,4-DIMETHYLPENTANAL	3-METHYLPENTANAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \text{C} \quad \text{C} \quad \text{C} \quad \text{C}=\text{C} \quad \text{C}-\text{C}-\text{C} \\ \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \end{array}$	(E+Z)-2-(1,2-DIMETHYLPROPYL)-4-METHYL-2-OCTENAL	UNK	3,4-DIMETHYLPENTANAL	2-METHYLPENTANAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \text{C} \quad \text{C} \quad \text{C} \quad \text{C}=\text{C} \quad \text{C}-\text{C}-\text{C} \\ \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \end{array}$	(E+Z)-2-(1,2-DIMETHYLPROPYL)-2-ETHYL-2-HEPTENAL	UNK	3,4-DIMETHYLPENTANAL	2-ETHYLPENTANAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \text{C} \quad \text{C} \quad \text{C} \quad \text{C}=\text{C} \quad \text{C}-\text{C}-\text{C} \\ \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \end{array}$	(E+Z)-2-PENTYL-4,6-DIMETHYL-2-HEPTENAL	UNK	HEPTENAL	2,4-DIMETHYLPENTANAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \text{C} \quad \text{C} \quad \text{C} \quad \text{C}=\text{C} \quad \text{C}-\text{C}-\text{C} \\ \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \end{array}$	(E+Z)-2-3-METHYLBUTYL-4,6-DIMETHYL-2-HEPTENAL	UNK	5-METHYLPENTANAL	2,4-DIMETHYLPENTANAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \text{C} \quad \text{C} \quad \text{C} \quad \text{C}=\text{C} \quad \text{C}-\text{C}-\text{C} \\ \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \end{array}$	(E+Z)-2-(1,2-DIMETHYLPROPYL)-4,6-DIMETHYL-2-HEPTENAL	UNK	3,4-DIMETHYLPENTANAL	2,4-DIMETHYLPENTANAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \text{C} \quad \text{C} \quad \text{C} \quad \text{C}=\text{C} \quad \text{C}-\text{C}-\text{C} \\ \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \end{array}$	(E+Z)-2-(1,2-DIMETHYLPROPYL)-4,6-DIMETHYL-2-HEPTENAL	UNK	3-METHYLPENTANAL	2-4-DIMETHYLPENTANAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \text{C} \quad \text{C} \quad \text{C} \quad \text{C}=\text{C} \quad \text{C}-\text{C}-\text{C} \\ \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \quad \text{C}^* \end{array}$	(E+Z)-2-(1-METHYLBUTYL)-6-DIMETHYL-2-HEPTENAL	TOTAL	93.1	

CARBANION AND ACCEPTOR COMPOUNDS PRESENT THE TWO ALDENES THAT WERE USED IN ALDOIS TO PREPARE THE TETRADEC-2-ENALS LISTED.

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EXAMPLE 18

This experiment and Example 19 illustrates the utility of the invention for mixtures of olefins. Hydroformylation of Dimate® hexenes and butenes was carried out in an autoclave with agitation. The 5 autoclave was charged with 2.61 g of dicobalt octacarbonyl(catalyst precursor), 270.2 g of Dimate® hexenes and 180.9 g butenes and 3000 psi gauge (20.786 KPa) of 1:1 CO and H₂. The autoclave was heated to 10 120°C and held for two hours. The autoclave and contents were cooled rapidly and the product removed under argon. The product was analyzed chromatographically as reported in Table 6.

TABLE 6

15	<u>Dimate® hexene, butene-2 Hydroformylation Products</u>	<u>Area %</u>
	<u>Compound</u>	
	Butene-2	1.5
	Dimate hexenes	9.6
20	2,2-dimethylpentanal	0.5
	2,4-dimethylpentanal	2.1
	2-methylbutanal	10.0
	Pentanal	23.6
	2-ethyl-3-methylbutanal	0.4
25	2-ethylpentanal	3.3
	3,4-dimethylpenntanal	0.5
	2-methylhexanal	3.2
	3-methylhexanal	18.5
	5-methylhexanal	12.5

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Heptanal	9.3
High Boilers (including heptyl alcohols and Formates)	5.0
The percent of aldehyde production without 5 2-substitution was over 76%.	

EXAMPLE 19

An aldehyde mixture representative of that from the oxo reaction of Dimate® hexenes and butene as described in Example 18 was reacted in an aldol condensation. A stirred autoclave was charged with 313 ml methanol, 0.8M NaOH, and 110 psi (758.4 KPa) gauge argon. The system was heated to 100°C and the mixture of aldehydes, 313 ml, was pressured into the autoclave with argon. The reaction was run an additional one hour and thirty minutes and the system rapidly cooled. The product was removed and the upper phase and lower phases were separated. The upper phase contained 0.5% unreacted pentanals, 31.7% unreacted heptanals, 25.6% decenals, 27.1% 20 2-dodecenals, 6.3% 2-tetradecenals, 1.7% pentanals, 6.3% heptanals and 0.8% high boilers. The skeletal structures of the 2-decenals and the 2-dodecenals are given in TABLE 7. The enals can readily be converted to alcohols and ethoxyl, sulfate or ethoxylsulfate 25 derivatives.

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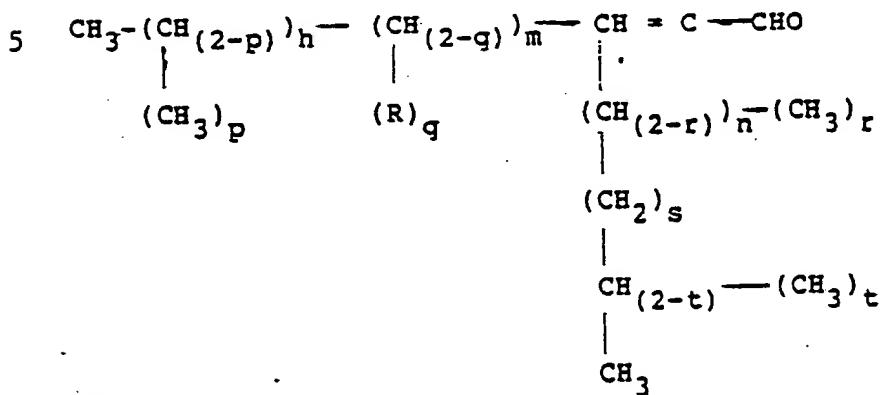
TABLE 7
 2-DECENALS AND 2-DODECENALS FROM ALDOL OF
 PENTANAL DIMATE HEXENES HEPTANALS

STRUCTURE	COMPOUND	NAME
C-C-C-C-C=C-C-C-C-C		(E+Z)-2-Propyl-2-Heptenal
2-DODECENALS		
C-C-C-C-C-C-C=C-C-C-C-C		(E+Z)-2-propyl-2-nonenal
C-C-C-C-C-C-C=C-C-C-C-C-C		(E+Z)-2-Propyl-7-Methyl-2-Octenal
C-C-C-C-C-C-C=C-C-C-C-C-C		(E+Z)-2-Propyl-5-Methyl-2-Octenal
C-C-C-C-C-C=C-C-C-C-C-C-C		(E+Z)-2-(1-Methylbutyl)-2-Heptenal
C-C-C-C-C=C-C-C-C-C-C-C-C		(E+Z)-2-(3-methylbutyl)-2-Heptenal
C-C-C-C-C=C-C-C-C-C-C-C-C		(E+Z)-2-Pentyl-2-Heptenal
C-C-C-C-C=C-C-C-C-C-C-C-C		(E+Z)-2-(1,2-Dimethylpropyl)-2-Heptenal
C-C-C-C-C=C-C-C-C-C-C-C-C		(E+Z)-2-Propyl-4,5-Dimethyl-2-Heptenal
C-C-C-C-C=C-C-C-C-C-C-C-C		(E+Z)-2-Propyl-4-Methyl-2-Octenal
C-C-C-C-C=C-C-C-C-C-C-C-C		(E+Z)-2-Propyl-4-Ethyl-2-Octenal

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The structure of most of the product is represented by the following formula:

Structure of C₁₀ to C₁₄ Enals from Aldol of Pentanal with heptanals from Oxo of Dimate Hexenes.



Wherein:

R = methyl or ethyl; h and m = 0, 1, 3, 4 and 5;
 h and s = 0, 1, 2; p, q, r and t = 0 or 1;
 m + h can never = more than 5

10 q and p = 0 when m + h = 5 or
 when m + l = 3

l = 3 and m = 1 when R = methyl and q = 1 and
 p = 0;

q = 0 and p = 1 when l = 2 and m = 2 or
 when l = 1 and m = 3;

15 n + s can never = more than 3
 r and t = 0 when n + s = 3 or
 when n + s = 1;

s = 0 when n, r and t = 1 or

20 when n = 2 and r = 0 and t = 1;

s = 1 when n = 1 and t and r = 0 or 1 but

only one of t and r can be 1;

s = 2 when t = 1 and n = 0;

h = 2 and m = 1 when R = ethyl and q = 1 and p = 0



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EXAMPLE 20

The autoclave of Example 1 was charged with 0.25 g carbonylbis (triphenylphosphine) rhodium (I) chloride, 11.78 g triphenylphosphine, 46.20 g 5 3-methylbutene-1 and 300 psi (2068.4 KPa) gauge 1:1 CO and H₂. The autoclave was heated to 130°C and held at this temperature for one hour and 35 minutes. The autoclave was cooled and the product removed under a 10 blanket of argon and analyzed. The product is 83% 4-methylpentanal with 7% 3-methylpentanal and 7% unreacted 3-methylbutene-1 starting material.

EXAMPLE 21

Hydroformylation was carried out on 2-methylbutene-1 with rhodium catalyst in accord with 15 the procedure of Exammple 20 at 100°C for an eight hour fifteen minute period. A 49% conversion was obtained with 98% selectivity to 3-methylpentanal.

EXAMPLE 22

An aldehyde mixture representative of that 20 from the oxo reactions as described above in Examples 20 and 21 was reacted in an aldol condensation. An autoclave was charged with 50 ml 0.8M NaOH, 50 ml methanol, and 10 psig (68.9 KPa) argon. The autoclave was heated to 100°C with agitation set at 1500 rpm and 25 the mixture of 23.08 g 3-methylpentanal and 23.11 g 4-methylpentanal, 50 ml total, were pressured into the autoclave with argon. The reaction was run for an additional hour and the system rapidly cooled. The product was removed and the upper and lower phases 30 were separated. The upper phase contained 2% methanol, 4.0% unreacted 3-methylpentanal, 2.6% unreacted 4-methylpentanal, 0.4% hexanols, 0.6% high boilers with 92.4% as dodecenals. The skeletal structures of the dodecenals are the same as the first 35 four structures of dodecenals listed in Table 8 which



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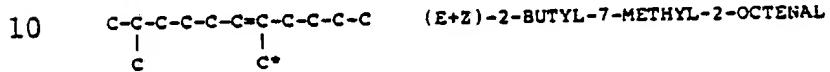
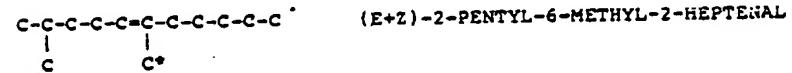
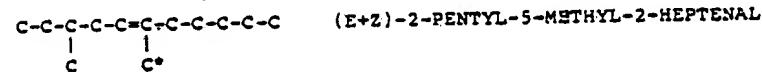
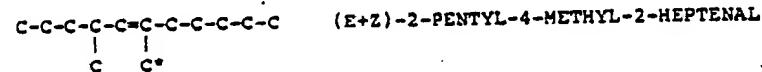
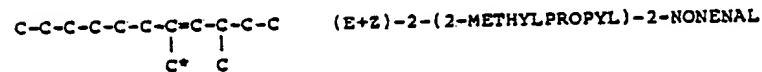
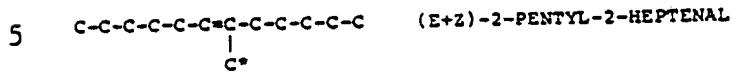
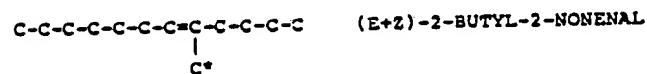
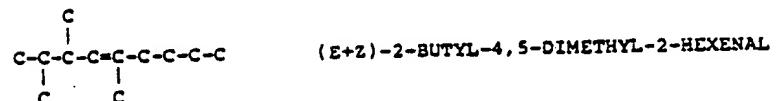
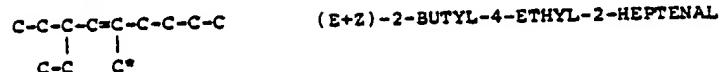
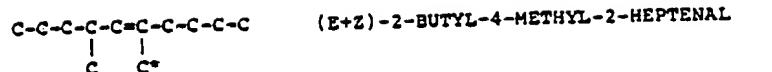
details skeletal structures of a mixed pentene stream with Dimate[®] hexenes, if hydroformylated with cobalt catalyst and aldol condensed as in Examples 17 and 18.

TABLE 6
2-DODECENALS AND 2-TRIDEcenals FROM Oxo AND ALDOL OF
PENTENES AND DIMATE⁵ HEXENES

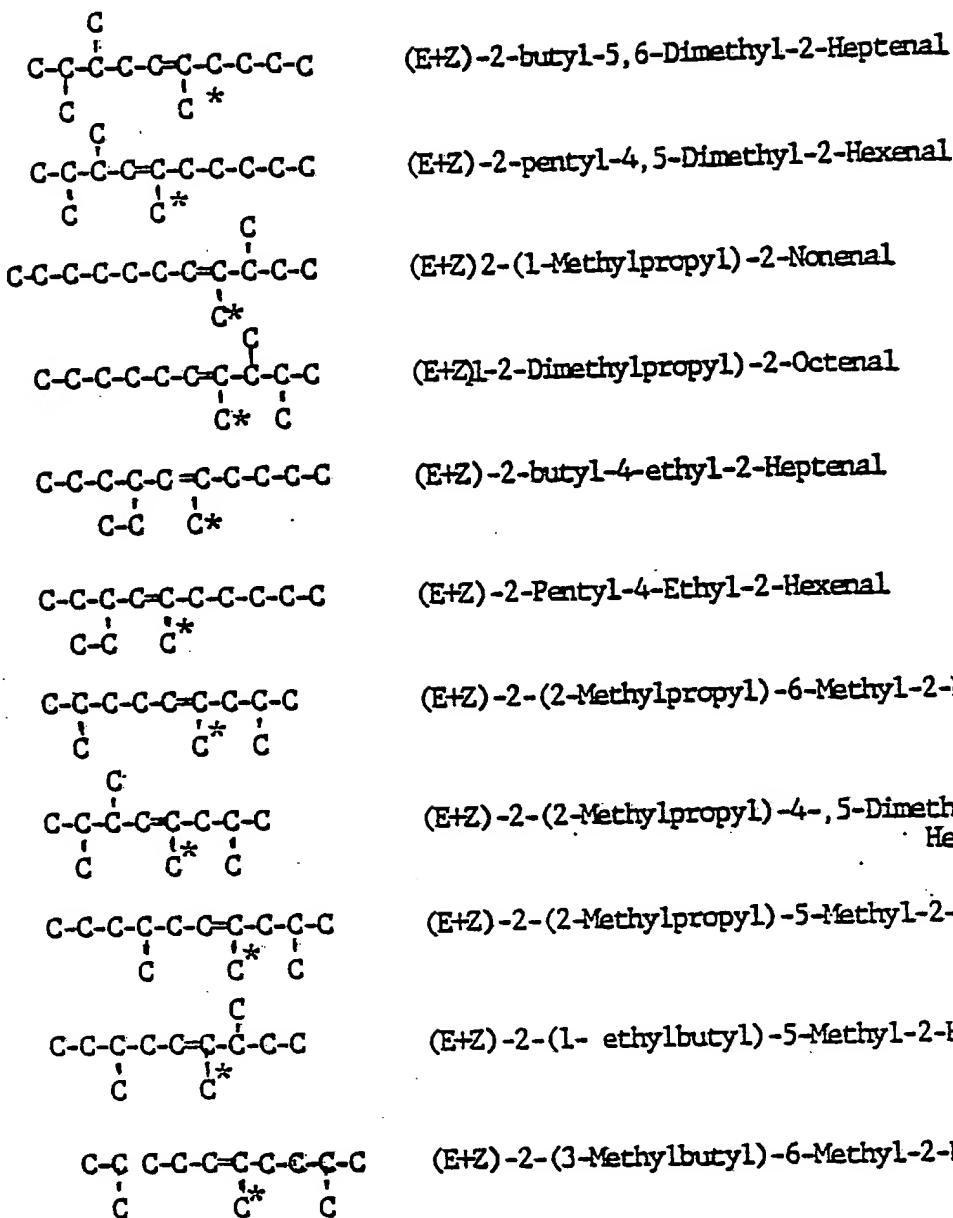
STRUCTURE	COMPOUND	NAME
		10
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C & C^* & C & & & \\ \end{array} $	(E+Z)-2-(2-METHYLPROPYL)-6-METHYL-2-HEPTENAL	
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C & C^* & C & & & \\ \end{array} $	(E+Z)-2-(2-METHYLPROPYL)-5-METHYL-2-HEPTENAL	
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C & & C^* & & & \\ \end{array} $	(E+Z)-2-(1-METHYLPROPYL)-6-METHYL-2-HEPTENAL	
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C & C^* & & & & \\ \end{array} $	(E+Z)-2-(1-METHYLPROPYL)-5-METHYL-2-HEPTENAL	
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C^* & & & & & \\ \end{array} $	(E+Z)-2-BUTYL-2-OCTENAL	
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C & C^* & & & & \\ \end{array} $	(E+Z)-2-BUTYL-6-METHYL-2-HEPTENAL	
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C & C^* & & & & \\ \end{array} $	(E+Z)-2-BUTYL-5-METHYL-2-HEPTENAL	
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C & C^* & & & & \\ \end{array} $	(E+Z)-2-BUTYL-4,5-DIMETHYL-2-HEXENAL	
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C^* & & & & & \\ \end{array} $	(E+Z)-2-(1-METHYLPROPYL)-2-OCTENAL	15
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C^* & & C & & & \\ \end{array} $	(E+Z)-2-(2-METHYLPROPYL)-2-OCTENAL	
$ \begin{array}{ccccccc} & C & & C & & C & \\ C-C-C-C-C=C & & & & & & \\ & C & C^* & C & & & \\ \end{array} $	(E+Z)-2-(1-METHYLPROPYL)-4,5-DIMETHYL-2-HEXENAL	



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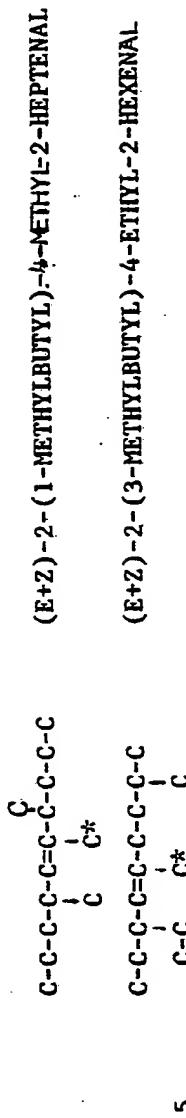
		$(E+Z)-2-(3\text{-METHYLBUTYL})-4,5\text{-DIMETHYL-2-HEPTENAL}$
		$C-C-C-C=C-C-C-C$ C C C C C C C C
5		$(E+Z)-2-(1\text{-METHYLPROPYL})-4,5\text{-DIMETHYL-2-HEPTENAL}$
		$C-C-C-C-C=C-C-C-C$ C C C C C C C C
10		$(E+Z)-2-(1\text{-METHYLPROPYL})-7\text{-METHYL-2-OCTENAL}$
		$C-C-C-C-C=C-C-C-C$ C C C C C C C C
15		$(E+Z)-2-(3\text{-METHYLBUTYL})-6\text{-METHYL-2-HEPTENAL}$
		$C-C-C-C-C=C-C-C-C$ C C C C C C C C
		$(E+Z)-2-(1\text{-METHYLBUTYL})-4\text{-ETHYL-2-HEXENAL}$
		$C-C-C-C-C=C-C-C-C$ C C C C C C C C

20

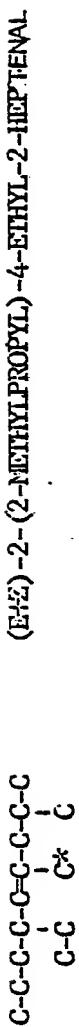
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$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-2-(1,2-\text{DIMETHYLPHENYL})-7-\text{METHYL}-2-\text{HEXENAL$
$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-2-(1,1-\text{DIMETHYLPHENYL})-5-\text{METHYL}-2-\text{HEXENAL$
$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-2-(1,1-LINOLEYLPHENYL)-4-\text{METHYL}-2-\text{HEPTENAL$
$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-2-(1,2-\text{DIMETHYLPHENYL})-2-\text{METHYL}-2-\text{HEXENAL$
$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-2-(2-\text{METHYLPHENYL})-6-\text{METHYL}-2-\text{OCTENAL$
$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-2-(1-\text{METHYLBUTYL})-4,4-\text{DIMETHYL}-2-\text{HEPTENAL$
$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-2-(1,2-\text{DIMETHYLPHENYL})-5-\text{METHYL}-2-\text{HEXENAL$
$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-1-(2-\text{METHYLPHENYL})-2-\text{OCTENAL$
$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-2-\text{ISOPROPYL}-5-\text{METHYL}-2-\text{OCTENAL$
$ \begin{array}{c} & \text{C} \\ & \\ \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ & & \\ \text{C} & \text{C} & \text{C} \end{array} $	$(E+Z)-2-(1-\text{METHYLPHENYL})-1-\text{OCTENAL$

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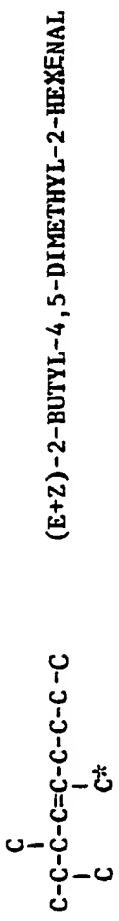
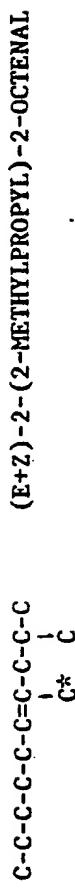
-70-

$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \qquad \qquad \\ \text{C} \qquad \text{C} \qquad \text{C} \\ \\ \text{C} \end{array}$	(E+Z)-2-(3-METHYLUTYI)-5-HEPTENAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \qquad \qquad \\ \text{C} \qquad \text{C} \qquad \text{C} \\ \\ \text{C} \end{array}$	(E+Z)-2-(1,2-DIMETHYLPHENYL)-6-HEPTENAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \qquad \\ \text{C} \qquad \text{C} \\ \\ \text{C} \end{array}$	(E+Z)-2-(1-METHYLUTYI)-4,5-DIMETHYL-2-HEPTENAL
$\begin{array}{c} \text{C} \qquad \text{C} \\ \qquad \\ \text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C} \\ \qquad \qquad \\ \text{C} \qquad \text{C} \qquad \text{C} \end{array}$	(E+Z)-2-(1,7-DIMETHYLPHENYL)-4,5-DIMETHYL-2-HEPTENAL
$\begin{array}{c} \text{C} \\ \\ \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C} \\ \qquad \\ \text{C} \qquad \text{C} \end{array}$	(E+Z)-2-(2-METHYLPHENYL)-4-METHYL-2-CYCLOHEPTENAL
$\begin{array}{c} \text{C} \\ \\ \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C} \\ \qquad \\ \text{C} \qquad \text{C} \end{array}$	(E+Z)-2-(1-METHYLPHENYL)-4-PHENYL-2-OCTENAL
$\begin{array}{c} \text{C} \\ \\ \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C} \\ \qquad \\ \text{C} \qquad \text{C} \end{array}$	(E+Z)-2-METHYL-4-METHYL-2-OCTENAL
$\begin{array}{c} \text{C} \\ \\ \text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C} \\ \qquad \qquad \\ \text{C} \qquad \text{C} \qquad \text{C} \end{array}$	(E+Z)-2-(1,2-DIMETHYLPHENYL)-4-METHYL-2-CYCLOHEPTENAL

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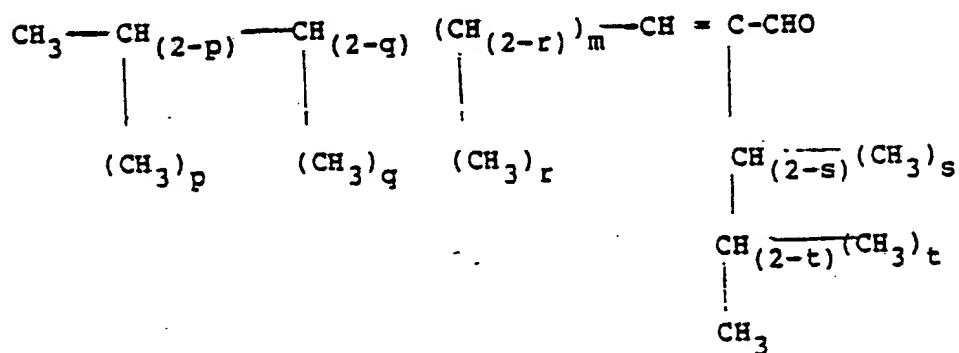
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Similarly, such oxo-aldol reactions of a predominant component of isoamylene streams, 2-methyl-butene-2, would produce an enal mixture with most components represented by the structure:

5 Structures of Dodec-2-enals from Oxo-Aldol of 2-Methylbutene-2



Wherein:

10 $m = 0$ or 1

$p, q, r, s, t = 0$ or 1 but only one of s and t can be 1 ;

$m = 1$ when $r = 0$ and only one of p and q can be 1 ;

$m = 0$ when p and $q = 1$

15 -----

At the oxo stage of the reaction the main aldehydes produced are 3-methylpentanal-1 and 4-methylpentanal-1 with possibly up to 10% 2,3-dimethylbutanal-1.

The aldol products are mainly 5-methyl-2-

20 (1-methylpropyl)hept-2-enal-1, 6-methyl-2-

(2-methylpropyl)-hept-2-enal-1, 6-methyl-

2(1-methylpropyl)hept-2-enal-1 and 5-methyl-2

(2-methylpropyl)hept-2-enal-1, and two C_{10} aldehydes in very small amounts.

25

EXAMPLE 23

The unsaturated aldehydes representative of



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product from oxo and aldol reactions of 2-methylbutene-1 and 3-methylbutene-1 as produced in Example 22 were subjected to hydrogenation. An autoclave was charged with 3.74 g of 45% + 5% cobalt 5 on Kieselghur, 37.27 g of the aldol condensation product, 23.16 g methanol and hydrogen to 1500 psi (10,644 KPa) gauge. The autoclave was heated to 160°C. The pressure and temperature were maintained for seventeen hours. The catalyst was filtered off 10 and washed with methanol to remove any residual alcohols. The dodecanols were distilled at 25 mm Hg and collected from 137 to 139°C.

EXAMPLE 24

15 This example illustrates the utility of the invention for synthesis of C₁₁ and C₁₄ unsaturated aldehydes.

An aldehyde mixture representative of that from the oxo reaction of Dimate® hexenes, as described above in Example 1 was reacted in an aldol 20 condensation along with normal butanal. A stirred autoclave was charged with 600 ml 0.8M NaOH, 1160 ml methanol and 20 psig (137.8 KPa) argon. The autoclave was heated to 100°C and a mixture of 323.2 g heptanals from oxo of Dimate® hexenes and 212.9 g butanal, 25 total volume 600 ml was pressured into the autoclave with argon. The reaction was run for an additional hour and the system rapidly cooled. The product was removed and the upper and lower phases were separated. The upper phase contained 2.7% methanol, 5.1% 30 unreacted heptanals, 13.7% (Z+E)-2-ethyl-hexenal, 56.0% 2-undecenals, 18.4% 2-tetradecenals, 0.5% heptanols and 3.5% high boilers. The mole ratio of the butanal to heptanals is 1.04:1, the mole ratio of the product aldehydes, 2-ethyl-hexenal, 2-undecenals 35 and 2-tetradecenals is 0.84 to 2.51 to 0.65



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respectively. The 2-undecenals were separated from the other products by reduced pressure distillation at 40 mm Hg. The 2-undecenals were collected in the temperature range of 119° to 132°C. The skeletal structures of the undecenals are given in Table 9.



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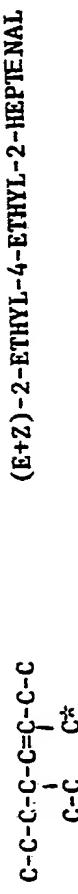
TABLE 9
2-UNDECENALS FROM ALDOL OF HEPTANALS AND BUTANAL

STRUCTURE	COMPOUND NAME
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C} \\ \\ \text{C} \end{array}$	(E+Z)-2-ETHYL-2-NONENAL
5 $\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C} \\ \\ \text{C} \end{array}$	(E+Z)-2-ETHYL-7-METHYL-2-OCTENAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C} \\ \\ \text{C} \end{array}$	(E+Z)-2-ETHYL-5-METHYL-2-OCTENAL
10 $\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C}-\text{C} \\ \\ \text{C} \end{array}$	(E+Z)-2-(1-METHYLBUTYL)-2-HEXENAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C}-\text{C} \\ \\ \text{C} \end{array}$	(E+Z)-2-(3-METHYLBUTYL)-2-HEXENAL
$\begin{array}{c} \text{C}-\text{C}-\text{C}-\text{C}=\text{C}-\text{C}-\text{C}-\text{C}-\text{C} \\ \\ \text{C} \end{array}$	(E+Z)-2-PENTYL-2-HEXENAL

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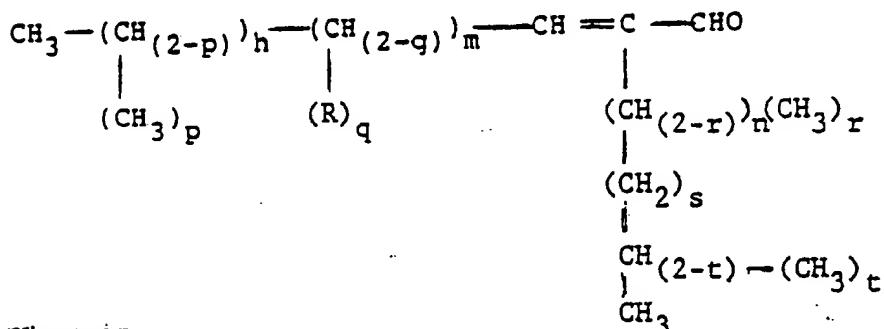
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The 2-undecenals and 2-tetradecenals in the product which are suitable for preparing detergent hydrophobes, can be represented by the following formula:

5 Structure of C₈, C₁₁ and C₁₄ Enals from Aldol of
Butanal
and Mixed Heptanals



Wherein:

10 R = methyl or ethyl; h and n = 0, 1, 2, 3, 4, and 5;
 n and s = 0, 1, 2; p, q, r and t = 0 or 1;
 q and p = 0 when m + k = 5 or when m + k = 2
 n + s can never be greater than 2
 m + h can never be greater than 5
 15 h = 3 and m = 1 when R = methyl and q = 1 and p = 0;
 h = 2 and m = 1 when R = ethyl and q = 1 and p = 0 or
 when q = 0 and p = 1;
 h = 1 and m = 3 when p = 1 and q = 0;
 r and t = 0 when n + s = 3 or when n + s = 0;
 20 s = 0 when n, r and t = 1 or when n = 2, r = 0 and t = 1;
 s = 1 when n = 1 and t and r = 0 or 1 but only one
 of t and r can be 1;
 s = 2 when t = 1 and n = 0



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EXAMPLE 25

This example illustrates the utility of the invention for a single hexene isomer, 4-methylpentene-1. The hydroformylation was performed in a 300 ml autoclave with agitation. The autoclave was charged with 0.29 g carbonyltris(triphenylphosphine)-rhodium hydride, 33.0 g triphenylphosphine, 100.0 g 4-methylpentene-1 and 200 psi (200 KPa) gauge pressure with 9 parts hydrogen to 1 part CO. The autoclave was heated to 100°C. and the pressure brought to 300 psi (2068.4 KPa) gauge using feed gas in 1 to 1 ratio hydrogen to CO. The temperature and pressure were maintained for 2 hours and 45 minutes after which the autoclave and contents were cooled rapidly. The product was removed and analyzed chromatographically. The product consists of 5.65% hexenes (5.57% 4-methylpentene-1), 10.64% 2,4-dimethylpentanal, 68.84% 5-methylhexanal, and approx. 15% triphenylphosphine, by area percent.

An aldehyde mixture representative of the oxo reaction of 4-methylpentene-1, as described above was reacted in an aldol condensation. The reaction vessel was a 100 ml 3 neck round bottom flask equipped with pressure equilizing funnel, thermometer, spiral condensor, a mechanical stirrer and a nitrogen blanket, maintained by using a slow bleed of nitrogen and a mineral oil bubbler. The reactor was charged with 12 ml 0.8M NaOH, and 12 ml 2,5-hexanediol, as co-solvent. The system was heated to 80°C and the aldehydes, 12 ml, were added over a six minute period. The system was heated to 89-90°C and held at temperature for an additional 30 minutes. The system was cooled and the upper and lower phases were separated. The upper phase was analyzed chromatographically and contained 2.0%



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2,4-dimethylpentanal, 1.5% 5-methylhexanal, 5% tetradecenal isomers, 85% 7-methyl-2-(3-methylbutyl)-2-octenal, 2.5% 2,5-hexanediol and a balance of high boilers.

5

EXAMPLE 26

The unsaturated aldehydes representative of product from oxo and aldol reactions of 4 methylpentene-1 as in example 25 were subject to hydrogenation. The reaction was performed in a 300 ml autoclave which was charged with 0.90 g 45 + 5% cobalt on Kieselghur, 7.92 g unsaturated aldehydes, 50.02 g methanol (used to ensure proper mixing) and 1500 psi (10644 KPa) gauge hydrogen. The autoclave was heated to 150°C. The pressure and temperature were maintained for one hour. The catalyst was filtered off and washed with methanol to remove any residual alcohols. The 7-methyl-2-(3-methylbutyl)-2-octenal was distilled at 20 mm Hg and was collected from 158 to 159°C. This example combined with Example 28 illustrates the conversion of the unsaturated aldehydes to saturated carboxylic acids.

EXAMPLE 27

Hydrogenation to a saturated aldehyde from an unsaturated aldehyde mixture representative of that from the oxo and aldol of Dimate R hexenes as described in Examples 1 and 17, was carried out in a 300 ml autoclave with agitation. The autoclave was charged with 2.60 g 5% palladium on carbon, 44.81 g methanol (used to ensure proper agitation), 8.24 g tetrade-2-enals and 200 psig (1379 KPa) H₂. The autoclave was heated to 85°C and held for one hour. The autoclave was then heated to 125°C and held for 2 hours. The autoclave was cooled and an infrared spectrometric analysis confirmed the reduction of the olefinic bond and the presence of a strong carbonyl



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band at 1740 cm^{-1} . The untreated 2-tetradecenals have bands at 1698 cm^{-1} , strong, and 1643 cm^{-1} moderate, which are characteristic of unsaturated aldehydes.

5

EXAMPLE 28

This example, combined with Example 27 illustrates the conversion of the unsaturated aldehydes to saturated carboxylic acids.

Oxidation of a saturated aldehyde mixture 10 representative of that from the hydrogenation of the unsaturated aldehydes as described in Example 27 was carried out in a Fisher-Porter aerosol bomb, with agitation. The bomb was charged with 60 ml heptene, 8.2 g tetradecanals, and 100 psi (689.4 KPa) gauge 15 oxygen. Agitation was started and pressure was allowed to drop to 60 psi (413.7 KPa) gauge when the bomb would be re-pressured to 100 psi gauge. The reaction continued for 3 1/4 hours and the temperature varied from 23° to 30°C. The pressure was released 20 and system heated to 80°C and held for 2 hours. The product was analyzed chromatographically and by gas liquid chromatography - mass spectrometry to confirm that saturated acid was formed, and 98% of the aldehydes were converted.

25

EXAMPLE 29

Oxidation of an unsaturated aldehyde mixture representative of that from the oxo and aldol reactions described in Examples 1 and 17, was carried out in a Fisher-Porter aerosol bomb with agitation.

30 The bomb was charged with 60 ml heptane, 8.2 g tetradec-2-enal and 100 psi (689.4 KPa) gauge oxygen in a darkened area. Agitation was started and the pressure in the bomb was allowed to drop to 70 psi (482.63 KPa) gauge at which time the bomb was 35 repressurized to 100 psi (689.4 KPa) gauge. The

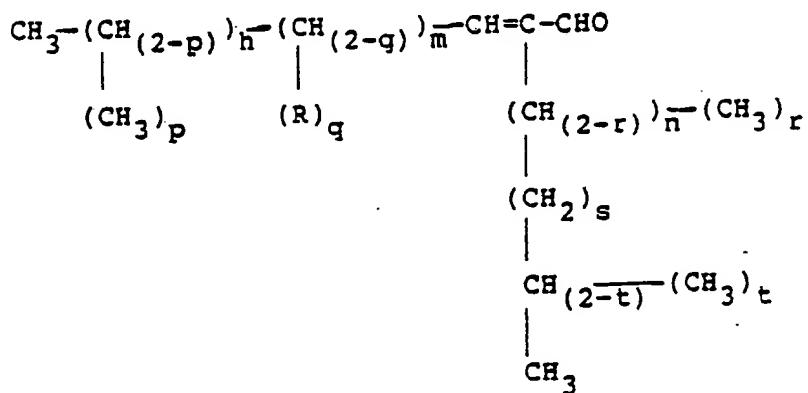


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reaction continued in this manner for four hours and the temperature varied from 20°C to 25°C. The pressure was released and the system heated to 80°C for 2 hours. The product was analyzed 5 chromatographically and by mass spectrometry and contained unsaturated tetradecanoic acids and 97% of the aldehydes were converted.

A mixture of amylenes and Dimate® hexenes can be subjected to oxo and aldol processes in accord 10 with the present invention. Thus, pentenes containing n-pentene-1, n-pentene-2, 3-methylbutene-1, 2-methylbutene-2, and 2-methylbutene-1 in admixture with Dimate® hexenes can be reacted in an oxo reaction, with the product treated in an aldol 15 reaction, in accord with procedures in Examples 18 and 19, and the resulting enals (see Table 8) can be represented by the following structure:

Structure of Dodec-2-enals and Tridec-2-enals
 20 from the, Oxo-Aldol of mixed pentenes
and Dimate Hexenes



Wherein:

R = methyl or ethyl; h and m = 0, 1, 2, 3, 4 and 5; n and s = 0, 1, 2; p, q, r and t = -0, 1;



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$m + h = 5$ or 4 , when p and $q = 0$;
 m and $q = 1$ and $p = 0$ when $h = 3, 2$ or 1 but
R can only be ethyl when $h = 1, 2$ and R
can only be methyl when $h = 2, 3$;

5 $h + m$ can never be greater than 5 ;
 $n + s$ can never be greater than 2 ;
 $q = 0$ and $p = 1$ and $h = 2$ when $m = 1$ or 0 ;
 $q = 0$ and $p = 1$ and $h = 1$ when $m = 3$ or 2 ;
 r and $t = 0$ when $n + s = 3$ or when $n = 0$ and $s = 2$;
10 $t = 1$ when $n = 0$ or 1 and $s = 1$ or 2 ;
 n and $t = 1$ when $s = 0$ and $r = 1$;

The aldehydes which are branched at the
2-position which react at a much slower rate in the
aldol reaction can be recycled to generate C_{11} - C_{16}
15 enals in several different ways. Thus these aldehydes
can be simply recycled to the aldol reaction until
their concentration is high enough that they are being
reacted in cross-aldol reactions at a rate essentially
the same as they are being generated in the
20 hydroformylation reaction. Another approach is to
allow these relatively unreactive aldehydes to proceed
through hydrogenation to give branched chain alcohols.
These alcohols can then be readily dehydrated to a
mixture of branched olefins. If these olefins are
25 recycled to the hydroformylation reaction they are
hydroformylated primarily to aldehydes without a
branch at the 2-position and hence they will undergo
the alcohol reaction and generate aldol products in
the desired C_{11} to C_{16} range. This can be illustrated
30 by considering the mixture of aldehydes generated by
the hydroformylation of Dimate ^R hexenes (see Table
2). The 2-methylhexanal, 2-ethylpentanal and
2,4-dimethylpentanal isomers pass through the aldol
reaction stage largely unreacted. On hydrogenation
35 they are converted to 2-methylhexanol, 2-ethylpentanol



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and 2,4-dimethylpentanol. These alcohols can be dehydrated by passage over heterogeneous catalysts at temperatures of 120-500°C. Catalysts should be chosen such that alcohol dehydration occurs readily but minimal skeletal rearrangement occurs. Suitable catalysts are various types of aluminas, thorias and metal phosphates (e. g. AlPO_4 , BPO_4 , $\text{Ca}_3(\text{PO}_4)_2$. (For further catalyst information see Journal of the American Chemical Society, 85, 2180 (1963) and Topics in Phosphorus Chemistry Vol. 10. page 285). As an example we can consider the recycle of 2-methylhexano. Over a dehydration catalyst it would give primarily 2-methylhexene-1 and 2-methylhexene 2. These olefins when charged to the hydroformylation reaction would give rise to primarily 3-methylheptanal with lesser amounts of 6-methylheptanal and minor amounts of other isomers. The 3-methylheptanal and 6-methylheptanal readily undergo aldol condensation and in conjunction with the C_7 aldehydes resulting directly from the hydroformylation of the Dimate^R hexenes will give rise to predominantly C_{15} unsaturated aldehydes with minor amounts of C_{16} unsaturated aldehydes. See Table 10 for specific names of the unsaturated aldehyde products. The described recycle involving dehydration and formation of aldolable aldehydes, permits handling smaller volumes of materials in the aldol reaction than is the case when the concentration of 2-branched aldehyde reactant is permitted to build up.



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TABLE 10
 2-PENTADECENALS AND 2-HEXADECENALS FROM OXO AND ALDOL OF
 DIMATE[®] HEXENES AND RECYCLED UNREACTIVE HEPTANALS

COMPOUND NAME

(E+Z)-2-(1-METHYL-PENTYL)-2-NONENAL
 (E+Z)-2-(1-METHYL-PENTYL)-7-METHYL-2-OCTENAL
 (E+Z)-2-(1-METHYL-PENTYL)-5-METHYL-2-OCTENAL
 (E+Z)-2-(1-METHYL-PENTYL)-5,6-DIMETHYL-2-HEPTENAL
 (E+Z)-2-PENTYL-5-METHYL-2-NONENAL
 (E+Z)-2-(1-METHYL-BUTYL)-5-METHYL-2-NCNENAL
 (E+Z)-2-(3-METHYL-BUTYL)-5-METHYL-2-NCNENAL
 (E+Z)-2-(1,2-DIMETHYL-PROPYL)-5-METHYL-2-NCNENAL
 (E+Z)-2-(1-METHYL-PENTYL)-4-METHYL-2-OCTENAL
 (E+Z)-2-(1-METHYL-PENTYL)-4-ETHYL-2-HEPTENAL
 (E+Z)-2-(4-METHYL-PENTYL)-2-NONENAL
 (E+Z)-2-(4-METHYL-PENTYL)-7-METHYL-2-OCTENAL
 (E+Z)-2-(4-METHYL-PENTYL)-5-METHYL-2-OCTENAL
 (E+Z)-2-(4-METHYL-PENTYL)-5,6-DIMETHYL-2-HEPTENAL
 (E+Z)-2-PENTYL-8-METHYL-2-NONENAL
 (E+Z)-2-(1-METHYL-BUTYL)-8-METHYL-2-NCNENAL
 (E+Z)-2-(3-METHYL-BUTYL)-8-METHYL-2-NCNENAL
 (E+Z)-2-(1,2-DIMETHYL-PROPYL)-8-METHYL-2-NCNENAL
 (E+Z)-2-(4-METHYL-PENTYL)-4-METHYL-2-OCTENAL
 (E+Z)-2-(4-METHYL-PENTYL)-4-ETHYL-2-HEPTENAL
 (E+Z)-2-(1-ETHYL-BUTYL)-2-NONENAL
 (E+Z)-2-(1-ETHYL-BUTYL)-7-METHYL-2-OCTENAL
 (E+Z)-2-(1-ETHYL-BUTYL)-5-METHYL-2-OCTENAL
 (E+Z)-2-(1-ETHYL-BUTYL)-5,6-DIMETHYL-2-HEPTENAL
 (E+Z)-2-PENTYL-5-ETHYL-2-OCTENAL
 (E+Z)-2-(1-METHYL-BUTYL)-5-ETHYL-2-OCTENAL
 (E+Z)-2-(3-METHYL-BUTYL)-5-ETHYL-2-OCTENAL
 (E+Z)-2-(1,2-DIMETHYL-PROPYL)-5-ETHYL-2-OCTENAL
 (E+Z)-2-(1-ETHYL-BUTYL)-4-METHYL-2-OCTENAL
 (E+Z)-2-(1-ETHYL-BUTYL)-4-ETHYL-2-HEPTENAL
 (E+Z)-2-PENTYL-4-ETHYL-6-METHYL-2-HEPTENAL
 (E+Z)-2-(1-METHYL-BUTYL)-4-ETHYL-6-METHYL-2-HEPTENAL
 (E+Z)-2-(3-METHYL-BUTYL)-4-ETHYL-6-METHYL-2-HEPTENAL
 (E+Z)-2-(1,2-DIMETHYL-PROPYL)-4-ETHYL-6-METHYL-2-HEPTENAL
 (E+Z)-2-PENTYL-4,7-DIMETHYL-2-OCTENAL
 (E+Z)-2-(1-METHYL-BUTYL)-4,7-DIMETHYL-2-OCTENAL
 (E+Z)-2-(3-METHYL-BUTYL)-4,7-DIMETHYL-2-OCTENAL
 (E+Z)-2-(1,2-DIMETHYL-PROPYL)-4,7-DIMETHYL-2-OCTENAL
 (E+Z)-2-PENTYL-4-ETHYL-5-METHYL-2-HEXENAL
 (E+Z)-2-(1-METHYL-BUTYL)-4-ETHYL-5-METHYL-2-HEXENAL
 (E+Z)-2-(3-METHYL-BUTYL)-4-ETHYL-5-METHYL-2-HEXENAL
 (E+Z)-2-(1,2-DIMETHYL-PROPYL)-4-ETHYL-5-METHYL-2-HEXENAL

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2-HEXADECENALS

(E+Z)-2-(4-METHYL-PENTYL)-7-METHYL-2-NONENAL
(E+Z)-2-(1-METHYL-PENTYL)-7-METHYL-2-NONENAL
(E+Z)-2-(1-ETHYL-BUTYL)-7-METHYL-2-NCNENAL
(E+Z)-2-(4-METHYL-PENTYL)-4,7-DIMETHYL-2-CCTENAL
(E+Z)-2-(1-METHYL-PENTYL)-4,7-DIMETHYL-2-OCTENAL
(E+Z)-2-(1-ETHYL-BUTYL)-4,7-DIMETHYL-2-OCTENAL
(E+Z)-2-(4-METHYL-PENTYL)-4-ETHYL-6-METHYL-2-HEPTENAL
(E+Z)-2-(1-METHYL-PENTYL)-4-ETHYL-6-METHYL-2-HEPTENAL
(E+Z)-2-(1-ETHYL-BUTYL)-4-ETHYL-6-METHYL-2-HEPTENAL
(E+Z)-2-(4-METHYL-PENTYL)-4-ETHYL-5-METHYL-2-HEPTENAL
(E+Z)-2-(1-METHYL-PENTYL)-4-ETHYL-5-METHYL-2-HEPTENAL
(E+Z)-2-(1-ETHYL-BUTYL)-4-ETHYL-5-METHYL-2-HEPTENAL
(E+Z)-2-(4-METHYL-PENTYL)-4-METHYL-2-NONEKAL
(E+Z)-2-(1-METHYL-PENTYL)-4-METHYL-2-NONENAL
(E+Z)-2-(1-ETHYL-BUTYL)-4-METHYL-2-NCNENAL
(E+Z)-2-(4-METHYL-PENTYL)-4-ETEYL-2-OCTENAL
(E+Z)-2-(1-METHYL-PENTYL)-4-ETEYL-2-OCTENAL
(E+Z)-2-(1-ETHYL-BUTYL)-4-ETHYL-2-CCTENAL

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Nonionic detergents prepared from the present alcohols by ethoxylation were tested for deteritive efficiency in comparison with reference compounds which were substantially normal alcohol ethoxylates, being Neodol [®] ethoxylates marketed by 5 Shell Chemical Company. The reference compound alcohols, produced by ethylene oligomerization, are composed of designated percentages of normal alcohols, generally in the range of 70 to 80%, and the remainder 10 of isomeric 2-alkyl (predominantly 2-methyl) primary alcohols. The tests employed are recognized tests in which a fabric soiled with synthetic sebum/airborne particulate is washed and the results measured by 15 Rd, change in reflectance by Gardner XL-23 Color Difference meter. Tests were conducted with a polyester-cotton blend fabric containing 65% polyester, and with a cotton broadcloth fabric. Three different ethoxylates of the C₁₄ alcohol mixture of 20 the present invention (designated C₁₄ Aldol) were used, having 6.0, 8.7 and 10.5 ethoxylate groups on the average. The reference compounds for comparison were Neodol 45 ethoxylates having 7 and 13 ethoxyl groups, and Neodol 25 ethoxylates having 7 and 12.5 ethoxyl groups. An individual oxo-alcohol C₁₄ alcohol 25 of the present invention was also tested 2-(1-methylbutyl)-5-methyloctanol. It was prepared by aldol of 3-methyl-heptanol, and is designated as C₁₄ High Vicinal, because of presence of adjacent branches 30 in its structure. The results are reported in Tables 11-a, 11-b, 11-c, 11-d, and 11-e. The tables include results with the detergent including sodium tripolyphosphate (STP) builder, sodium carbonate builder, or no builder, and a washing temperature of 120°F in most instances, but 75°F in Table 11-d.



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Tests were reported at different amounts of water hardness as set forth. The sodium tripolyphosphate built detergent, for example, had a composition of 10% of the nonionic (surfactant), 24% sodium tripolyphosphate, 12% R.U. Sebacate (as is), 53% sodium sulfate, and 1% sodium carboxyethyl cellulose, while the one with no builder was similar, with the STP replaced by additional sodium sulfate. The detergent was used in concentration 0.15%, for anionic concentration of 150 ppm. Water hardness was from a 3/2 atom ratio of calcium and magnesium ions, with concentration calculated as parts per million by weight of calcium carbonate.

15

Table 11aSynthetic Sebum Detergency

120°F, Cotton Δ Rd

30% Na₂CO₃ Builder System

20

	Water					
	<u>Sample</u>	<u>Hardness</u>	<u>0</u>	<u>50 ppm</u>	<u>100 ppm</u>	<u>200 ppm</u>
25	Neodol	45-7	32.5	31.5	29.6	27.2
	"	45-13	32.3	31.1	29.2	23.3
30	Neodol	25-7	31.8	29.9	28.8	23.9
	"	25-12.5	33.1	30.7	27.6	20.2
35	C ₁₄ Aldol	6.0	32.7	29.4	26.0	19.8
	"	8.7	32.6	30.8	27.0	24.0
	"	10.5	31.5	31.1	28.0	26.2
40	C ₁₄ High Vicinal	5.5	20.8	20.6	18.8	11.4
		9.5	19.5	18.9	19.4	14.8
		13.3	15.5	17.0	15.8	11.1



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Table 11bSynthetic Sebum Detergency

120° F, cotton, Δ Rd

0% Builder System

5	Water					
	<u>Sample</u>	<u>Hardness</u>	0	50 ppm	100 ppm	200 ppm
	Neodol		31.5	29.5	26.1	24.0
	"		31.4	31.3	29.5	27.8
	Neodol	25-7	31.7	29.4	26.1	24.8
10	"	25-12.5	30.1	31.1	29.7	25.9
	C ₁₄ Aldol-	6.0	30.4	26.0	20.7	16.8
	"	8.7	30.2	30.6	24.5	22.4
	"	10.5	30.4	30.9	26.3	24.7
	C ₁₄ High					
15	Vicinal	5.5	32.7	18.9	21.0	13.9
		9.5	32.1	20.1	20.3	17.4
		13.3	32.6	17.5	16.1	14.3



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Table 11-cSynthetic Sebum Detergency

120°F, Cotton, △ Rd
24% STP Builder System

5	<u>Sample</u>	Water Hardness	0 50 ppm 100 ppm 200 ppm		
			0	50 ppm	100 ppm
	Neodol	45-7	32.2	34.2	35.2
	"	45-13	31.5	34.0	35.0
10	Neodol	25-7	32.2	34.4	34.5
	"	25-12.5	31.7	33.5	34.5
	C ₁₄ Aldol	6.0	31.4	34.4	34.5
	"	8.7	30.8	33.5	35.7
	"	10.5	31.4	34.3	34.8
15	C ₁₄ High Vicinal	5.5	32.8	35.3	36.6
		9.5	32.8	33.5	35.8
		13.3	32.7	35.1	34.5
					22.4
					25.3
					24.9

Table 11-dSynthetic Sebum Detergency

20 75°F, PE/cotton, △ Rd
24% STP Builder System

<u>Sample</u>	Water Hardness	0 50 ppm 100 ppm 200 ppm		
		0	50 ppm	100 ppm
Neodol	45-7	18.3	17.7	15.7
				14.0



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	"	45-13	16.7	16.1	15.1	11.9
	Neodol	25-7	18.8	19.8	18.0	14.3
	"	25-1.25	16.5	17.2	14.5	13.7
5	C_{14} Aldol-	6.0	19.8	17.7	19.3	9.4
	"	8.7	19.3	18.2	20.0	15.0
	"	10.5	17.5	18.1	16.6	14.0
10	C_{14} High Vicinal	5.5	20.8	20.6	18.8	11.4
		9.5	19.5	18.9	19.4	14.8
		13.3	15.5	17.0	15.8	11.1

Table 11-eSynthetic Sebum Detergency

120°F, PE/cotton, Δ Rd
24% STP Builder System

15		Water				
		Sample	Hardness	0	50 ppm	100 ppm
	Neodol	45-7	19.8	19.7	18.7	16.2
	"	45-13	18.6	17.8	16.8	12.8
20	Neodol	25-7	19.9	19.6	20.4	16.7
	"	25-12.5	18.1	17.8	16.6	13.4
	C_{14} Aldol	6.0	19.8	19.0	18.6	7.9
	"	8.7	19.9	21.7	21.1	18.9
	"	10.5	20.1	18.8	19.6	17.6
25	C_{14} High Vicinal	5.5	21.2	18.9	21.0	13.9
		9.5	19.4	20.1	20.3	17.4



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13.3 18.4 17.5 16.1 14

It can be seen that the ethoxylates of the present alcohol mixture are in general comparable in the tests of detergative efficiency to the established reference ethoxylates. It can be noted that in some respects the present compounds give better results, as for example with the 8.7 ethoxylate in Tables 11-d and 5-e involving tests with polyester/cotton.

The C_{14} mixed alcohol product of the present invention, in ethoxylate form, was tested for biodegradability, in comparison to commercial detergents. The test procedure used was the semi-continuous activated sludge test which determines dissolved organic carbon as the measure of biodegradation. The results are reported in Table 12 below. The aldol-alcohol (C_{14}) was the alcohol mixture as described in Table 4 above, which has been ethoxylated to have an average of 10.5 ethoxyl groups. It was compared to a Neodol 25-12, which is an ethoxylate of an approximately linear alcohol in the 12 to 15 carbon range, with an average of 12 ethoxyl groups. The two LAS reference compounds are linear alkyl benzene sulfonates of a type utilized commercially. The test-measures the removal of dissolved organic carbon (DOC), and is a measure of complete degradation of the compounds.



Table 12

Test Compound	Mean % DOC Removal ¹ (95% Confidence Limits)		
	2nd through 5th Week of Tests		6th through 8th Week of Tests
	24-Hour Cycle	72-Hour Cycle	24-Hour Cycle
(16 data points)	(3 data points)	(12 data points)	(3 data points)
Neodol 25-12 Reference Compound	90 ± 6	98 ± 7	97 ± 2
A230 LAS Reference Compound	83 ± 7	92 ± 11	89 ± 3
Dodecen-1, Derive d LAS Reference Compound	81 ± 13	82 ± 59	89 ± 6
Aldol Alcohol (C ₁₄) Ethoxylate (10.5 EO)	77 ± 6	92 ± 16	88 ± 5
			100 ± 1

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¹ Mean % DOC Removal = (DOC in Feed - Net DOC in Effluent) X 100/DOC in Feed

Net DOC in Effluent = Test Chamber DOC - Control Chamber DOC



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While the degradation of the aldol product was not as fast as that of the linear alcohol product (Neodol 25-12), it was still comparable to the LAS compounds which are suitable for commercial use. Also, the 5 essentially complete removal in the 72-hour cycles indicates there are no resistant fragments.

Further biodegradation tests were carried out utilizing the semi-continuous activated sludge test, and comparing a C₁₄ aldol isomeric mixture ethoxylate with an ethoxylate of a high vicinal alcohol, i.e., 3-(1-methyl-butyl)-5-methylheptanol. The 10 24-hour cycle results for weeks 4, 5 and 6 are reported in Table 7. While the high vicinal alcohol degrades somewhat more slowly, its rate is still 15 fairly close to that of the isomeric mixture which was shown to be comparable to commercial materials in Table 6. Moreover, results with the isomeric mixture indicate that good biodegradation can be obtained even with a substantial amount of the high vicinal alcohol 20 present. The results in Table 7 provide a comparison of the materials tested, but may not be directly comparable to other tests, because of the low ethoxyl content which may have affected solubility, or possibly unexplained test variations.



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Table 13Semi-Continuous Activated Sludge Test

(% DOC. Removal - 24 hr)

			<u>Week 4</u>	<u>Week 5</u>	<u>Week 6</u>	
5	1.	Neodol ^R 25-7 Reference compound	88	38	97	13
	2.	C ₁₄ + 6.0 Ethylene Oxide (Dimersol Hexene)	61	13	65	31
	3.	High Vicinal Alcohol + 6.3 Ethylene Oxide (from 3-methylhexanal)	66	44	51	39
10					63	12



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EXAMPLE 30

The C₁₄ ethoxylate derivatives described above and evaluated were prepared by known ethoxylation procedures. A 100 ml glass reactor was employed with thermometer, stirrer, and provision for ethylene oxide addition. A mercury-filled U-tube manometer was employed, and about 700 mm pressure was maintained during the reaction. Ethylene oxide was charged from a small bomb, with the amount charged determined by weight difference. The weight of the detergent alcohol charge was calculated to have the desired mole ratio to ethylene oxide. The reaction temperature was maintained at 165°C by a controlled heating mantle. As catalyst, solid KOH was charged at 1% by weight of alcohol used. Thus C₁₄ aldol alcohol product, like that of Table 3, was reacted in separate reactions with ethylene oxide in mole ratio of 5.3/1 and 8.9/1 EO/alcohol by weight. The ethylene oxide addition rate was controlled by a valve and required about 2 to 2.5 hours for complete addition as determined by the manometer. The temperature was reduced to 125°C. and the catalyst was neutralized by addition of H₃PO₄ by syringe through a rubber septum. The amount of the acid in ml of 85% H₃PO₄ was 0.7 times the KOH weight in grams. The product was held for 15 minutes at 125°C following the acid addition, and was then filtered while hot through a layer of Celite® filter aid on a fritted filter to remove phosphate salts. NMR analysis showed ethylene oxide content similar to that charged in the reactions, being 5.6 ethoxyl units and 8.7 ethoxyl units respectively. A similar procedure but with a higher EO/alcohol charge was used to prepare an ethoxylate of the same alcohol mixture, but with 10.5 ethoxyl units



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per alcohol molecule. The procedure was also used to prepare the C₁₄ high vicinal alcohol evaluated herein, using weight charges of EO/alcohol of 5.9, 8.8 and 12.1 to obtain products with 5.5, 9.5 and 13.3 ethoxyl 5 units as shown by NMR.

EXAMPLE 31

Sulfate derivatives of the alcohols or ethoxylates were prepared by sulfation with chlorosulfonic acid. Both an alcohol mixture, and an 10 ethoxylated alcohol mixture were sulfated in this manner. The alcohol mixture was from oxo, aldol and hydrogenation reactions starting with Dimate R hexenes, as described hereinabove. The ethoxylate was prepared in accord with the above procedure with an 15 ethylene oxide charge calculated to provide three ethoxylate units. The chlorosulfonic acid was provided on a 1.05 to 1 mole ratio to alcohol or ethoxylate. In separation reactions, the alcohol and ethoxylate were placed in flasks and cooled in an ice 20 bath while acid was added dropwise. HCl vapors were then drawn off by intermittent vacuum. When HCl evolution stopped, the sulfated products were transferred to beakers containing solutions of 90% isopropanol and 10% water, by weight. The sulfate 25 derivatives were 20% by weight of their respective solutions. A 50% by weight solution of NaOH in water was added with agitation to pH 8.0. Salts solids were removed by filtration. The solutions of sulfate derivatives were then evaporated to dryness on a 30 rotary evaporator. The products were then in the form of a sodium sulfate of the C₁₄ alcohol mixture, and a sodium sulfate of the ethoxylated C₁₄ alcohol mixture containing a nominal 3.0 ethoxyl units. Ethanol was added to the products to form concentrated solutions, 35 along with water to aid solubility, forming a 55.6%



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solution of the alcohol sulfate, and a 50.2% solution of the sulfate of the ethoxylated alcohol. The solutions are a convenient form for preparing samples for evaluation of detergency, biodegradation and 5 similar properties. In view of the results reported herein for ethoxyl derivatives, the sulfate derivatives are expected to be similarly good.

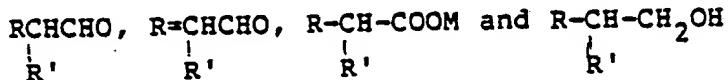
The detergent hydrophobes of the present invention are also of interest in the form of 10 synthetic fatty acids. The long chain aldehydes as produced herein, can readily be converted to acids by simple oxidation of the aldehyde group, as in an air oxidation. The acids retain the branched structure of the aldehyde, and can have the unsaturation at the 15 2-position as in the enals, or such unsaturation can be hydrogenated, or otherwise reduced, to provide saturated acids. Such acids can be converted to sodium, potassium or other alkali metal soaps or other detergents having anionic or hydrophilic groups 20 attached to the acid, and are expected to have suitable lathering and other detergent properties. It will be noted that the acids, which can be termed as synthetic fatty acids, have the same structure as the alcohols. Since the alcohols are suitable as 25 detergent hydrophobes, having suitable detergent and biodegradation properties, the corresponding acids are expected to have suitable detergent and biodegradation properties.



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CLAIMS:

1. Mixtures containing a number of isomeric compounds selected from those of structures:



5 having from 11 to 16 carbon atoms, in which R has 6 to 8 carbon atoms, and R' has from 3 to 6 carbon atoms, with most of the compounds having additional methyl or ethyl branches, and in which M is hydrogen or a salt-forming cation.

10 2. A mixture of isomeric alcohols having from 11 to 16 carbon atoms and characterized as alkanols of 6 to 10 carbon atoms with an alkyl group containing 2 to 6 carbon atoms substituted on the 2-position thereof, and with additional branching in 15 most of the isomers, with most of the additional branches being methyl groups, and further characterized as being liquid at ambient temperatures and having hydrocarbon hydrophobe moieties making the mixture suitable for formation of effective detergents 20 therefrom, and further characterized by suitable biodegradability.

3. A mixture of compounds as defined in Claim 2, but in which the alcohols have been converted to ethoxyl derivatives with an average of about 3 to 12 ethoxyl units per alcohol unit.

25 4. The mixture of Claim 3 in which the ethoxyl units are in the range on the average of 6 to 12 units per alcohol unit.

5. A mixture of compounds as in claim 2 in which the alcohols have been converted to sulfate 30 derivatives.



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6. A mixture of compounds as in Claim 2 in which the alcohol has been converted to ethoxysulfate derivatives.

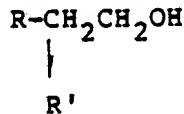
7. A mixture of isomeric enals having from 5 11 to 16 carbon atoms and characterized as 2-alkenals of 6 to 10 carbon atoms with an alkyl group containing 2 to 6 carbon atoms substituted in the 2-position thereof, and with additional branching in most of the isomers with most of the additional branches being 10 methyl groups, and further characterized as being liquid at ambient temperatures and having hydrocarbon hydrophobe moieties making the mixture suitable for formation of effective detergents therefrom, and further characterized by suitable biodegradability.

15 8. The mixture of Claim 7 in which the aldehydes on the average have more than 2 branches in addition to the 2-branch.

9. The mixture of Claim 7 in which aldehydes of different carbon numbers are present.

20 10. A mixture of C₁₄ isomeric alcohols which are characterized as nine-carbon alkanol with a five-carbon alkyl group substituted on the 2-position thereof, and with additional branching in most of the isomers, with most of the additional branches being 25 methyl groups, and further characterized as liquid at ambient temperature and having varying hydrocarbon hydrophobe moieties making the mixture suitable for formation of effective detergents therefrom, and further characterized by suitable biodegradability.

30 11. The alcohol mixture of Claim 10 in which the alcohols are characterized by the structure:



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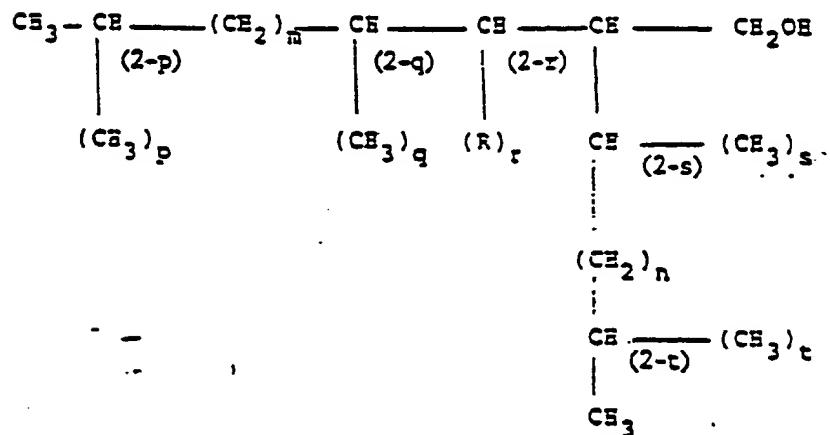
in which R is an alkyl group with 7 carbon atoms and R' is an alkyl group with 5 carbon atoms and in which in at least about 80% of the alcohols, R' is selected from n-pentyl, 3-methylbutyl and 1-methylbutyl, and R 5 is selected from n-heptyl, 3-methylhexyl, 5-methylhexyl, 2-methylhexyl and 2-ethylpentyl groups.

12. The alcohols of Claim 10 in which about 40 to 60% by weight of the alcohols are 2(3-methylbutyl)-5-methyloctanol, 2-10 (1-methylbutyl)-5-methyloctanol and 2-pentyl-5-methyloctanol.

13. The alcohols of Claim 12 in which about 65% to about 80% by weight of the alcohols are comprised of the alcohols named in Claim 12 along with 15 2-(3-methylbutyl)-7-methyloctanol, 2-pentynonanol, 2-pentyl-7-methyloctanol and 2-(3-methylbutyl)-nonanol.

14. The alcohols of Claim 12 further characterized in that about 20 to about 35% of the 20 alcohol have adjacent branches in their structure, but there are substantially no alcohols with gem substituents and therefore substantially no quaternary carbon.

15. An alcohol mixture of Claim 10 having 25 most of the alcohols represented by the structure:



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wherein:

R = methyl or ethyl;
p, q and r = 0 or 1; but only one of p, q and r
can be 1;
m = 3, when p, q and r = 0;
m = 2, when r = 1 and R = methyl;
m = 1, when r = 1 and R = ethyl;
n = 2, when s and t = 0;
n = 1, when s or t = 1;
s and t = 0 or 1, but only one of s and t can be 1.

16. Detergent composition comprising
ethoxyl, sulfate or ethoxylsulfate derivatives of the
alcohols in the mixture of Claim 10.

5 17. Detergent compositions comprising
ethoxyl, sulfate or ethoxylsulfate derivatives of the
alcohols of Claim 14.

18. As a new compound, a C₁₄ alcohol
suitable for use in formation of nonionic detergents
10 and in liquid form, designated as 2-(3-methylbutyl)
-5-methyl-octanol.

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19. As a new compound, a C₁₄ alcohol suitable for use in formation of nonionic detergents and in liquid form, designated as 2-(1-methylbutyl)-5-methyloctanol.

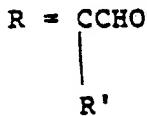
5 20. As a new compound, a C₁₄ alcohol suitable for use in formation of nonionic detergents and in liquid form, designated as 2-pentyl-5-methyloctanol.

10 21. As a new compound, a C₁₄ alcohol suitable for use in formation of nonionic detergents and in liquid form, designated as 2-(3-methylbutyl)-7-methyloctanol.

15 22. The alcohol mixture of Claim 10 further characterized as having alcohols present substantially in amounts as set forth in Table 4.

20 23. A mixture of C₁₄ isomeric enals which are characterized as nine-carbon 2-alkenal with a five-carbon alkyl group substituted on the 2-position thereof, and with additional branching in most of the isomers, with most of the additional branches being methyl groups, and further characterized as liquid at ambient temperature and having varying hydrocarbon hydrophobe moieties making the mixture suitable for formation of effective detergents therefrom, and 25 further characterized by suitable biodegradability.

24. A mixture of enals characterized by the structure:



30 in which R is an alkylidene group with 7 carbon atoms and R' is an alkyl group with 5 carbon atoms and in which in at least about 80% of the compound, R' is selected from n-pentyl, 3-methylbutyl and

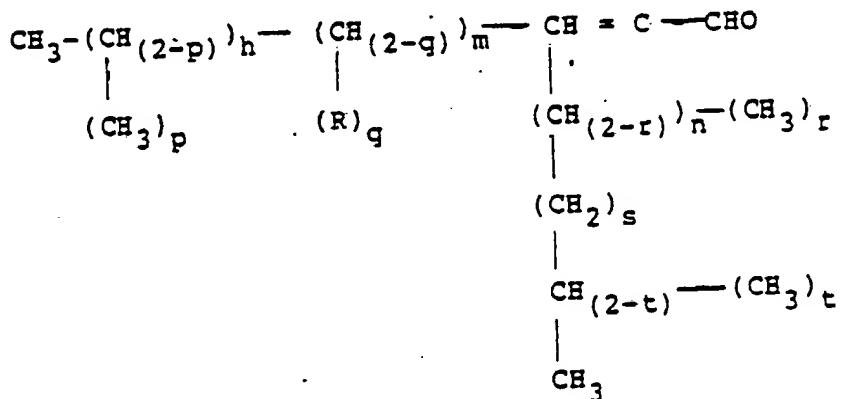


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1-methylbutyl, and R is selected from n-pentyl,
 3-methylbutyl and 1-methylbutyl, and R is selected
 from n-heptylidine, 3-methylhexylidene, 5
 methylhexylidene, 2-methylhexylidene and
 5 2-ethylpentylidene groups.

25. A mixture of enals characterized by
 structure as set forth in Claim 15, but with the
 $-\text{CH}_2\text{OH}$ group replaced by $-\text{CHO}$, and with a 2, 3 double
 bond.

10 26. A mixture of isomeric enals in accord
 with Claim 7 and further characterized as having most
 of the aldehydes in the mixture represented by the
 structure set forth below, and their saturated
 aldehyde, alcohol, ethoxylated alcohol, and
 15 ethoxylated alcohol su fa e derivatives:



Wherein:

R= methyl or ethyl; h and m = 0, 1, 3, 4 and 5;
 h and s = 0, 1, 2; p, q, r and t = 0 or 1;
 m + h can never = more than 5
 q and p = 0 when m + h = 5 or
 when m + 1 = 3
 1 = 3 and m = 1 when R= methyl and q = 1 and
 p=0;

q = 0 and p = 1 when 1 = 2 and m = 2 or
 when 1 = 1 and m = 3;

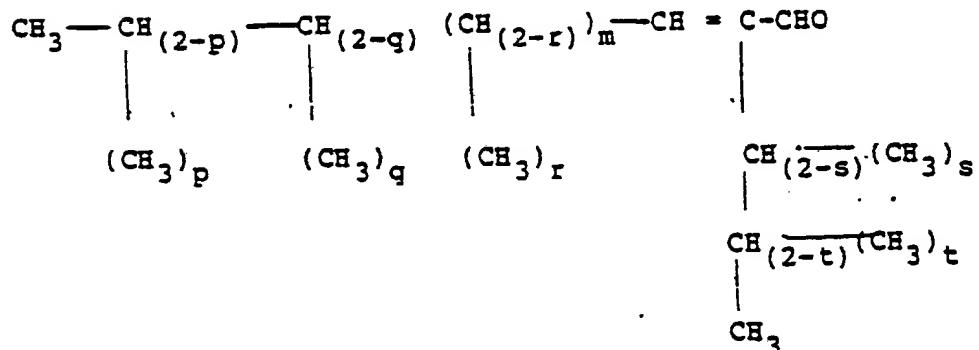
n + s can never = more than 3
 r and t = 0 when n + s = 3 or



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when $n + s = 1$;
 $s = 0$ when n, r and $t = 1$ or
when $n = 2$ and $r = 0$ and $t = 1$;
 $s = 1$ when $n = 1$ and t and $r = 0$ or 1 but
only one of t and r can be 1;
 $s = 2$ when $t = 1$ and $n = 0$;
 $h = 2$ and $m = 1$ when $R =$ ethyl and $q = 1$ and $p = 0$

27. A mixture of isomeric enals in accord
with Claim 7 and further characterized as having most
of the aldehydes in the mixture represented by the
5 structure set forth below, and their saturated
aldehyde, alcohol, ethoxylated alcohol, and
ethoxylated alcohol sulfate derivates:



Wherein:

$m = 0$ or 1

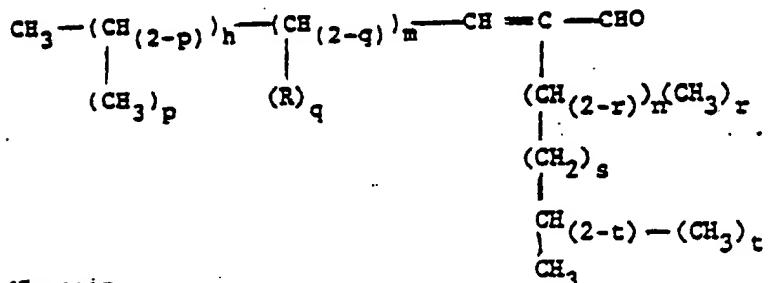
$p, q, r, s, t = 0$ or 1 but only one of s and t can be 1;

$m = 1$ when $r = 0$ and only one of p and q can be 1;

$m = 0$ when p and $q = 1$



28. A mixture of 18 meric enals in accord with Claim 7 and further characterized as having most of the aldehydes in the mixture represented by the structure set forth below, and their saturated aldehyde, alcohol, ethoxylated alcohol, and ethoxylated alcohol sulfate derivatives:



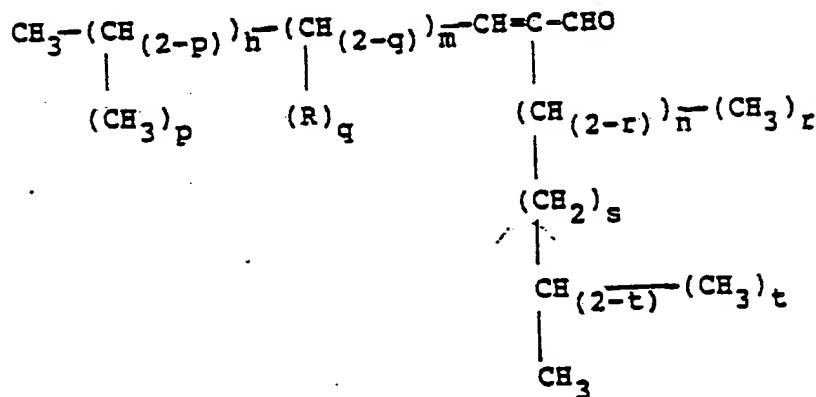
Wherein:



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$m + h = 5$ or 4 , when p and $q = 0$;
 m and $q = 1$ and $p = 0$ when $h = 3, 2$ or 1 but
 R can only be ethyl when $h = 1, 2$ and R
 $can only be methyl when h = 2, 3$;
 $h + m$ can never be greater than 5 ;
 $n + s$ can never be greater than 2 ;
 $q = 0$ and $p = 1$ and $h = 2$ when $m = 1$ or 0 ;
 $q = 0$ and $p = 1$ and $h = 1$ when $m = 3$ or 2 ;
 r and $t = 0$ when $n + s = 3$ or when $n = 0$ and $s = 2$;
 $t = 1$ when $n = 0$ or 1 and $s = 1$ or 2 ;
 n and $t = 1$ when $s = 0$ and $r = 1$;

29. A mixture of isomeric enals in accord with Claim 7 and further characterized as having most of the aldehydes in the mixture represented by the structure set forth below, and their saturated aldehyde, alcohol, ethoxylated alcohol, and ethoxylated alcohol sulfate derivatives:



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Wherein:

R = methyl or ethyl; h and m = 0, 1, 2, 3, 4 and 5;

n and s = 0, 1, 2; p, q, r and t = -0, 1;

m + h = 5 or 4, when p and q = 0;

m and q = 1 and p = 0 when h = 3, 2 or 1 but

R can only be ethyl when h = 1, 2 and R
can only be methyl when h = 2, 3;

h + m can never be greater than 5;

n + s can never be greater than 2;

q = 0 and p = 1 and h = 2 when m = 1 or 0;

q = 0 and p = 1 and 1 = 1 when m = 3 or 2;

r and t = 0 when n + s = 3 or when n = 0 and s = 2;

t = 1 when n = 0 or 1 and s = 1 or 2;

n and t = 1 when s = 0 and r = 1;



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30. The enals of Claim 23 in which about 40 to 60% by weight of the enals are 2(3-methylbutyl)-5-methyl-2-octenal, 2(1-methylbutyl)-5-methyl-2-octenal and 2-pentyl-5-methyl-2-octenal.

5 31. As a new compound, a C₁₄ enal suitable for conversion to compounds for use in detergents, and in liquid form, designated as 2(3-methylbutyl)-5-methyl-2-octenal.

10 32. As a new compound a C₁₄ enal suitable for conversion to compounds for use in detergents, and in liquid form, designated as 2(1-methylbutyl)-5-methyl-2-octenal.

15 33. A process of preparing detergent range aldehydes which comprises conducting oxo reactions in which olefins are reacted with hydrogen and carbon monoxide to provide aldehydes and conducting an aldol reaction of such aldehydes comprised of at least 50% alkanals with no branching at the 2-position and of carbon number so as to produce aldol product having mainly carbon numbers in the range of 11 to 16 carbon atoms, to cause good conversion to aldol product but with that produced from cross-aldol of 2-substituted aldehydes constituting no more than 20% of the product.

25 34. The process of Claim 33 in which product is obtained in enal form.

30 35. The process of Claim 33 in which the aldol reaction is conducted in aqueous alkali and a cosolvent is present which aids in improving contact of the aldehydes with the aqueous alkali.

36. The process of Claim 35 in which an hydroxyl compound is employed as cosolvent.

37. The process of Claim 33 in which the



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aldol product is hydrogenated and oxidized to obtain corresponding saturated carboxylic acids.

38. The process of Claim 33 in which the aldol product is hydrogenated to alcohol and therefrom 5 converted to ethoxyl, sulfate or ethyoxyl sulfate derivative.

39. The process of Claim 33 in which the oxo reaction is conducted with cobalt catalyst at 10 temperatures in the range of 80 to 150°C and pressures sufficient to maintain catalyst stability but not over 5000 psi (34,475K Pa), and the aldol reaction is conducted in aqueous alkaline medium at temperatures of about 90 to about 130°C.

40. The process of Claim 33 in which 15 aldehyde for the aldol reaction has content of alkanals with no 2-branching of about 60 to about 80% and the conversions of aldehyde to aldol product are in the range of about 75% to about 90%.

41. A process of preparing detergent range 20 aldehydes which comprises conducting an oxo reaction with hydrogen, carbon monoxide and olefins selected from those having 5 to 7 carbon atoms to obtain aldehydes having from 6 to 8 carbon atoms and comprised of at least 50% alkanals with no branching 25 at the 2-position and subjecting the aldehydes to an aldol reaction to cause good conversion to aldol product but with that produced from cross-aldol of 2-substituted aldehydes constituting no more than 20% of the product.

42. The process of Claim 41 in which the aldol reaction is conducted in aqueous alkali and a cosolvent is present which aids in improving contact 30 of the aldehydes with the aqueous alkali.

43. The process of Claim 42 in which an 35 hydroxyl compound is employed as cosolvent.



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44. The process of Claim 43 in which the cosolvent is methanol or an alkane diol.

45. The process of Claim 41 in which a C₅ olefin is reacted.

5 46. The process of preparing detergent range aldehydes with a high content of aldehyde in the C₁₁ to C₁₄ range which comprises conducting an aldol reaction of heptanals obtained from oxo reaction of branched hexenes from propylene dimerization, with 10 butanal and obtaining an aldol product with a high content of 2-undecenals.

47. The process of Claim 46 in which aldol product is hydrogenated to alcohol and ethoxylated to an ethoxyl containing detergent.

15 48. The process of Claim 46 in which aldol product is hydrogenated to alcohol and converted to a sulfate with detergent properties.

20 49. The process of preparing aldehydes which comprises conducting an oxo reaction with hydrogen carbon monoxide and a hexene mixture comprises mainly of methylpentenes with internal unsaturation with no more than 30% linear hexenes so as to obtain C₇ aldehydes with less than 30% 2-substituted aldehydes and subjecting the aldehydes 25 to aldol reaction to obtain aldol product with conversion of at least 70%.

30 50. A process for preparing detergent range aldehydes which are liquid and suitable for conversion to alcohols for use in formation of effective and biodegradable detergents, from olefin feedstock in an efficient process with good yields compared to other routes for producing detergent alcohols from olefins, which comprises conducting an oxo process with cobalt catalyst with a hexene mixture obtained by 35 dimerization of propylene and comprised mainly of



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methylpentenes to obtain aldehyde product with at least 75% or so of aldolable aldehyde, and subjecting the aldehydes to an aldol reaction with basic catalyst to obtain enals which are liquid at ambient

5 temperature and comprising C₁₄ enal isomers characterized as nine carbon enals with a five carbon alkyl substituent on the 2-position thereof and with additional, mostly methyl branching in most isomers.

10 51. The process of Claim 49 in which the aldol product is hydrogenated to alcohols, including hydrogenation of aldehydes which did not react at the aldehyde stage, separating the resulting C₇ alcohols and dehydrating same to C₇ olefins and cycling the olefins back to the oxo process to convert such

15 52. olefins to C₈ aldehydes which are then subjected to the aldol reaction and subsequent hydrogenation with aldol product, thereby producing an alcohol product comprised mainly of C₁₅ isomers corresponding to the C₁₄ enals, but with some C₁₄ and C₁₆ alcohols from 20 participation of C₈ aldehyde in the aldol reaction.

53. The process of Claim 49 in which the hexenes contain a substantial amount of 2-methyl-2-pentene from which the aldehyde product is composed of more than 85% aldolable product and the aldolable content of the aldehyde mixture is better than 75%.

30 54. The process of Claim 41 in which the hexenes include 4-methyl-pentene-1 produced by alkali catalyzed dimerization of propylene.

55. A process of preparing detergent range aldehydes which comprises conducting oxo reactions in which olefins are reacted with hydrogen and carbon



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monoxide to provide aldehydes and conducting an aldol reaction of such aldehydes comprised of at least 50% alkanals with no branching at the 2-position and of carbon number so as to produce aldol product having 5 mainly carbon numbers in the range of 11 to 16 carbon atoms, and with the slower reacting 2-branched aldehyde being recycled to the aldol reaction to generate a higher yield of the desired unsaturated aldehyde.



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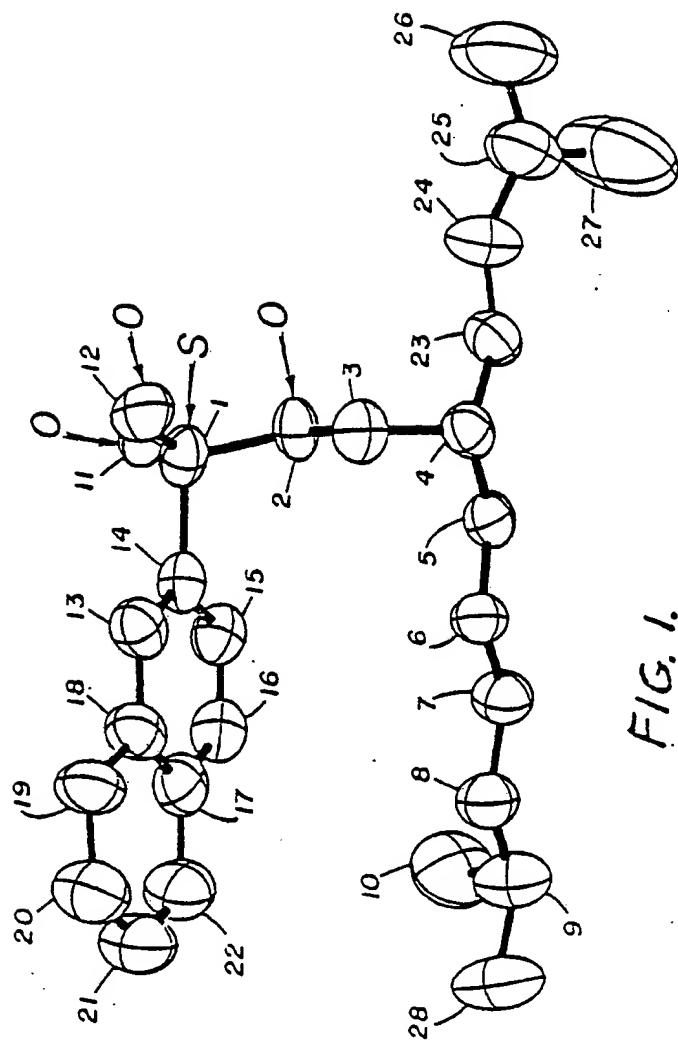
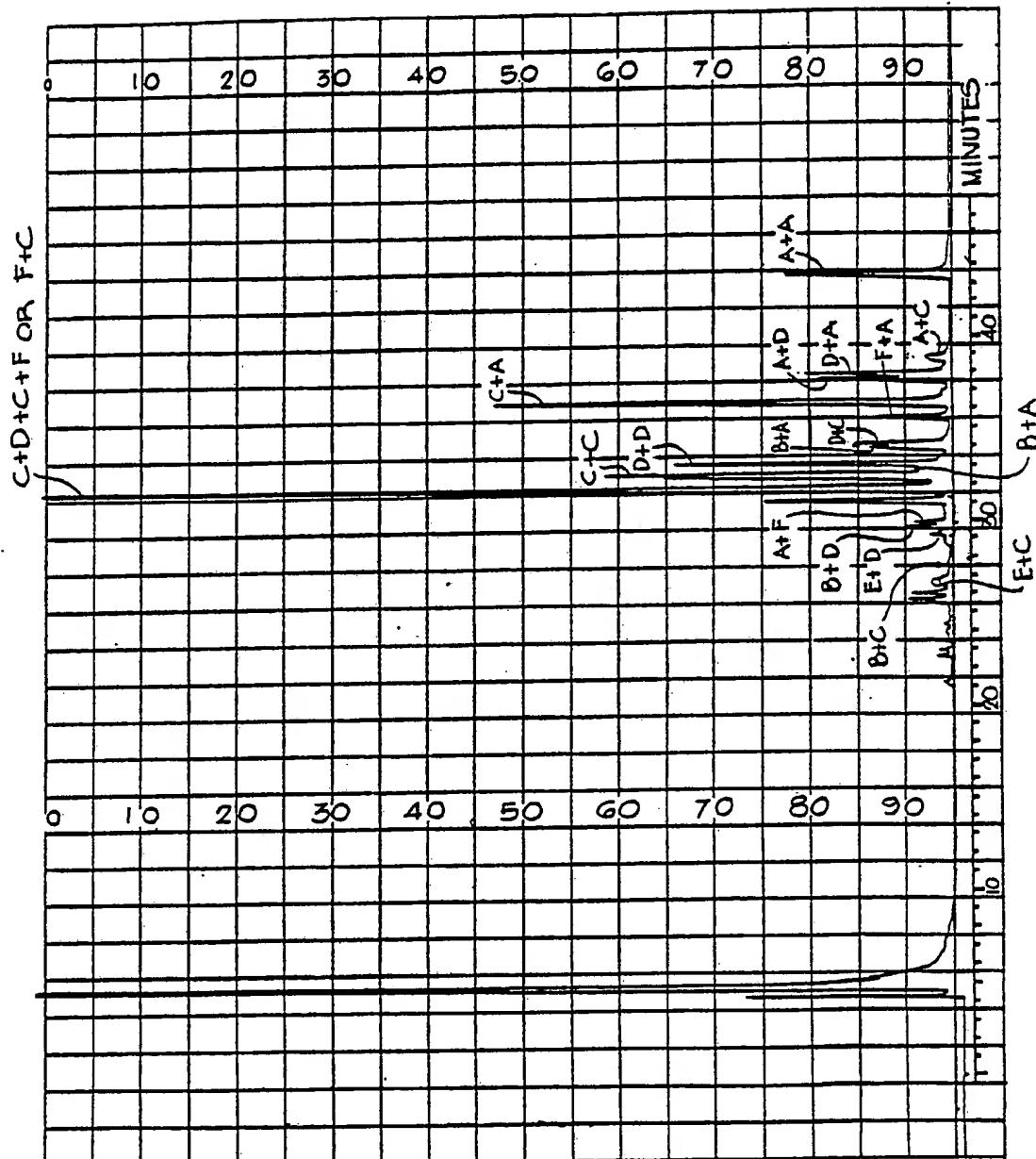


FIG. 1.

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60m x 0.25mm SP. 2100
 38 CSiH₆ 1x10⁻¹¹

60:1 SPLIT
 120 → 150°C @ 2°/min

0.1ml
 EFSC₁₄ ALL MIX
 +AE_{2C} +FYR

FIG.2.

SUBSTITUTE SHEET



INTERNATIONAL SEARCH REPORT

International Application No PCT/US83/01744

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all):

According to International Patent Classification (IPC) or in both National Classification and IPC
 INT. CL. 3 C07C 29/14, 45/00, 47/02, 47/20, 43/11, 51/235
 C11D 1/14, 1/29, 1/72

II. FIELDS SEARCHED

Minimum Documentation Searched⁴

Classification System	Classification Symbols
US	252/174.21, 550, 551 260/398.6 568/451, 463, 876, 878, 882, 909

Documentation Searched other than Minimum Documentation
 to the Extent that such Documents are Included in the Fields Searched⁵

NONE

III. DOCUMENTS CONSIDERED TO BE RELEVANT¹⁴

Category ⁶	Citation of Document, ¹⁵ with indication, where appropriate, of the relevant passages ¹⁷	Relevant to Claim No. ¹⁸
E, X	US, A 4,426,542 (BARKER) 17 January 1984	1-4, 6-36, 39-55
Y	US, A 4,032,578 (SAVINI) 28 June 1977	1, 2, 7-15, 18-36, 39-46, 49-55
X, Y	US, A 3,127,451 (BERKELEY) 31 March 1964	1, 2, 7-15, 18-36 39-46, 49-55
X, Y	US, A 3,119,876 (JAROS) 28 January 1964	1, 2, 7-15, 18-36 39-46, 49-55
X, Y	US, A 2,934,568 (BARKER) 26 April 1960	3, 4, 16, 17, 38, 47
X, Y	US, A 2,852,563 (HAGEMEYER) 16 September 1958	1, 2, 7-15, 18-37 39-46, 49-55
X, Y	US, A 2,655,525 (BANES) 13 October 1953	5, 16, 17, 38, 48
X, Y	US, A 2,654,772 (PAVLIC) 6 October 1953	5, 16, 17, 38, 48

* Special categories of cited documents:¹⁵

- "A" document defining the general state of the art which is not considered to be of particular relevance
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- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search¹⁹

29 FEBRUARY 1984

Date of Mailing of this International Search Report²⁰

02 MAR 1984

International Searching Authority²¹

RO/US

Signature of Authorized Officer²⁰

Dennis L. Albrecht

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